

SITE EVALUATION

80 LISTER AVENUE

**SUBMITTED TO
NEW JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION**

**PREPARED BY
DIAMOND SHAMROCK CHEMICALS COMPANY
IT CORPORATION
WOODWARD-CLYDE CONSULTANTS
ENVIRO-MEASURE, INC.**

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EXECUTIVE SUMMARY

The property located at 80 Lister Avenue in the Ironbound section of Newark, New Jersey is a former chemical manufacturing facility operated by a number of firms, including Diamond Shamrock Chemicals Company (Diamond Shamrock). An investigation conducted by the New Jersey Department of Environmental Protection (NJDEP) and the U.S. Environmental Protection Agency (EPA) confirmed in May 1983 the presence of dioxin at the 80 Lister Avenue property. Diamond Shamrock entered into a consent order agreement with NJDEP in March 1984, agreeing to secure the site, to take measures to minimize public exposure to contamination, and to undertake a site evaluation and a feasibility study of remedial alternatives. This report documents the site evaluation.

As part of the site evaluation, a Work Plan was prepared and submitted to NJDEP in June 1984. The Work Plan was modified in response to NJDEP comments and the amended Work Plan was approved in August 1984. A field investigation was subsequently conducted at the site between September and November 1984. Analytical laboratory testing has been performed on the various samples obtained at the site and the results are reported herein along with the results of other testing and other site characterization data.

Chemical manufacture at the 80 Lister Avenue site began in the early part of this century, but the period of most significance relative to the contamination observed at the site is from the end of World War II to the mid-1970's. During this time pesticides and phenoxy herbicides were the primary products manufactured, and dioxin occurs as a contaminant in some of these products.

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other 663 samples were analyzed for one or more of the following parameters as specified by the Work Plan:

- o 2,3,7,8-tetrachlorodibenzo-p-dioxin (2,3,7,8-TCDD or dioxin)
- o Volatile priority pollutants (38 compounds)
- o Semi-volatile priority pollutants (69 compounds)
- o Pesticides/herbicides (35 compounds)
- o Inorganic/classical parameters (15 compounds)
- o 2,3,7,8-tetrachlorodibenzofuran (2,3,7,8-TCDF)
- o Octachlorodibenzo-p-dioxin (OCDD)
- o Asbestos
- o Various ambient air/industrial hygiene parameters.

Results of the chemical analyses confirmed the presence of dioxin and a limited number of priority pollutant compounds on the site. Generally, the priority pollutant compounds detected with the greatest frequency and in the highest concentrations were those chemicals associated with previous site operations.

The site evaluation has established the following:

- o Dioxin is present both inside and outside all the site structures, with the highest level of contamination in the process and chemical manufacturing buildings. Because of the extent of contamination and materials of construction, decontamination is impractical.
- o Approximately 70 percent of the existing on-site drums are contaminated with dioxin.
- o All of the tanks, sewers, and sumps sampled are contaminated with dioxin.

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1.0 INTRODUCTION

This report presents the results of the site evaluation conducted at the 80 Lister Avenue site (hereinafter "the site"). The work has been performed for Diamond Shamrock Chemicals Company (Diamond Shamrock) by IT Corporation (ITC), and its subcontractors--Woodward-Clyde Consultants (WCC) and Enviro-Measure, Inc. (EMI). This report is being submitted to the New Jersey Department of Environmental Protection (NJDEP) in response to Administrative Consent Order I (ACO I) related to the site.

The report describes in detail the history of the site, the regional environmental setting, and all the activities associated with the site evaluation as defined in the 80 Lister Avenue Work Plan (Work Plan). Data associated with the field sampling and testing and analytical laboratory testing are presented, and these data are subsequently used to characterize the site with respect to the presence of 2,3,7,8-TCDD (dioxin) and the U.S. Environmental Protection Agency (EPA) priority pollutants plus 40 tentatively identified nonpriority pollutants in the volatile organic analysis, base/neutral, and acid fractions for each sample analyzed. Finally, the available information characterizing the site is evaluated to determine if an adequate data base exists for performing a feasibility study for remedial action. The following paragraphs describe the main sections of the report in more detail.

Section 2.0 presents the history of the site from its earliest known use as an industrial site to its current condition. Particular emphasis has been placed upon the period during which the hazardous materials known to be present were manufactured. A history of events leading to the site evaluation presented herein is also provided.

Section 3.0 presents the regional environmental setting of the site covering climate and meteorology, geology and landforms, surface and ground water hydrology, flora and fauna, and land usage. Specific

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An evaluation of the results of the investigation of the site is provided in Section 7.0. The data are carefully examined from the standpoint of providing an adequate definition of contamination of the items characterized in Section 6.0, particularly with regard to adequacy for performing a feasibility study. Areas of deficiency are clearly identified and a recommended program for obtaining any required additional data is presented.

References, tables, and figures for each section are provided at the end of that section. Tables and figures are numbered sequentially according to the section of the text in which they are first referenced. For example, the first two tables referenced in Subsection 2.6.2 would be 2.6.2-1 and 2.6.2-2. Appendices are presented sequentially at the end of the report and are identified by letters (Appendix A, Appendix B, etc.). A list of the material contained in each appendix is provided at the beginning of that appendix. The report Table of Contents is provided at the front of each of the volumes of the report.

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2.0 SITE HISTORY AND EXISTING CONDITIONS

2.1 SITE LOCATION

The 80 Lister Avenue site is located in the Ironbound section of Newark, New Jersey. The site occupies approximately 3.4 acres on the north side of Lister Avenue. It is nearly rectangular in shape, extending about 375 feet in an east-west direction and 405 feet north-south. The site is bounded on the north by the Passaic River, on the east by the former Sergeant Chemical Company site now owned by Diamond Shamrock, at the southeast corner by the Duralac Company property, and on the south and west by Sherwin-Williams Company property. Vehicular access to the site is via a common right-of-way shared with Duralac and which enters the southeast corner of the property. The property is formally described as Lots 58 and 59 in Block 2438 on the tax maps of Newark.

The location of the site within Newark and the Ironbound section is shown on the accompanying maps (Figures 2.1-1 and 2.1-2).

2.2 SITE HISTORY

2.2.1 History Prior to Purchase by Diamond Shamrock

Industrial development on the site is reported to date from the 1870's. Drawings from 1914, revised in 1922, show the site to be part of the Lister Agricultural Chemical Company property which extended for some distance along the Passaic River. This plant site also included most of the other nearby industrial sites.

It was during the period of ownership by Lister that the site reached its present dimensions following filling along the south shore of the Passaic River to form the northernmost 30 percent of the property (Figure 2.2.1-1). Much of the remainder of the site is also filled with the granular material reportedly used to fill the marsh land that existed in the natural state. Several buildings were on the site including the

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Ownership by Kolker ceased in March 1951 when the Kolker Chemical Works was acquired by Diamond Alkali Company (now Diamond Shamrock Chemicals Company).

2.2.2 History During Ownership by Diamond Alkali and Diamond Shamrock
(March 1951 to March 1971)

During this period the manufacture of several products was either transferred to other locations or discontinued, leaving the phenoxy herbicides as the only products of the plant. A major impetus for this change was an explosion in February 1960 which destroyed several plant processes--when rebuilt the plant only included processes for the manufacture of the phenoxy herbicides and their intermediates. The layout of the plant site prior to the explosion is shown in Figure 2.2.2-1. Modernization and expansions continued during the 1960's which saw total phenoxy capacity more than double to 15 million pounds per year. Principal events during Diamond Shamrock and subsequent ownership are listed in Table 2.2.2-1.

The changes started in 1955 with the transfer of Lindane manufacture to another location. Production of low γ -BHC continued until 1957 or 1958 when it also was relocated. The biggest change, however, was the transfer of DDT production, which was moved to Texas in late 1958 or early 1959. During the late 1950's several process changes were instituted to improve the operating efficiency of the plant. Among these was a change instituted around 1956 to the trichlorophenol (TCP) process to recycle trichloroanisole (TCA). A change in the handling of process effluent also occurred in 1956 with the installation of an industrial sewer connecting to the Passaic Valley Sewerage Commission (PVSC) Lister Avenue line. Following installation of that connection, most of the plant process wastes were discharged through the PVSC treatment plant.

An explosion in the TCP unit during February 1960 destroyed the large five-story building in which it and several other plant processes had been located. Following the explosion, a decision was made to limit

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- o 1963 - The 2,4-D acid process was rehabilitated. The roof was raised permitting installation of new ventilating ducts to carry process fumes to a new and larger caustic scrubber.
- o 1965 - The melt, washing, and drying process for the production of dry, flaked 2,4-D was installed, with a 40 percent increase in capacity. These changes also reduced personnel contact with the 2,4-D.
- o 1967 - The final plant expansion saw the construction of a new and larger 2,4-D unit and the conversion of the former 2,4-D unit to the manufacture of 2,4,5-T. The TCP purification process for dioxin removal via carbon filtration was added as part of this same expansion.

Operation at the plant continued until August 1969 when it was shut down. The production units were cleaned out as they were shut down, and between September and December the remaining raw materials and products were sold and shipped. The plant was listed for sale and remained idle throughout 1970 until it was purchased by Chemicaland Corporation in March 1971. It is noted that Chemicaland actually purchased the 1.8 acres and improvements owned by Diamond Shamrock, which then assigned rights to the 1.6 acres it had leased from Walter Ray Holding Company to Chemicaland.

2.2.3 History During Chemicaland and Subsequent Ownership (March 1971 to the Present)

Following purchase of the property by Chemicaland, equipment was installed for the manufacture of benzyl alcohol which was to be made and sold by Cloray NJ Corporation, an affiliate of Chemicaland. Production of benzyl alcohol was not profitable, so they attempted to expand their product line by manufacturing several specialty items and also by performing custom manufacture on a toll basis. These efforts were all unsuccessful and production ceased during the summer of 1973.

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Little is known of the use of the property by Marisol, but eventually this company started cleaning and clearing the site. Concerning the cleanup, it is known that:

- o The product left in the equipment when the plant was shut down on February 24, 1977, was removed and placed in drums, of which 570 remain on site today.
- o Some equipment known to be on the site following the shutdown was removed.
- o Warehouse space and tankage was leased to SCA Corporation which used it in conjunction with waste disposal operations at their neighboring plant. The date that SCA started to use the site is not exactly known, but was prior to the summer of 1982.

During the spring of 1983, SCA continued to lease and use a portion of the site, while Marisol was working to prepare the office building for occupancy. This was the situation in May 1983 when results of samples taken in April by the EPA showed high levels of dioxin on the site and NJDEP moved to control access to the property. On June 2, 1983, New Jersey Governor Kean issued Executive Order No. 40 which has guided control and cleanup activities since that date.

2.3 GENERAL PROCESS DESCRIPTION

The processes described in the following paragraphs relate to the manufacture of 2,4-D and 2,4,5-T and their intermediates at the time the plant was shut down by Diamond Shamrock. These processes are most relevant to the current state of contamination at the site.

2.3.1 MeTCP

MeTCP was made in a batch reaction by the alkaline hydrolysis with caustic soda of 1,2,4,5-tetrachlorobenzene (T₄CB) in the presence of methanol at temperatures and pressures near 165 degrees Centigrade and 350 psig, respectively. Methanol was distilled from the reaction mass

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2.3.5 2,4-D

Production of 2,4-D was carried out in batches by the alkaline condensation of the 2,4-DCP and MCA in the presence of caustic soda. The resulting sodium 2,4-D slurry was filtered and washed to remove unwanted sodium 2,6-D which was discarded. The sodium 2,4-D was then continuously acidified with sulfuric acid, washed to remove sodium sulfate, and dried in an evaporator. The resulting anhydrous, molten 2,4-D was stored for later flaking or for use in the preparation of the various 2,4-D esters or Dacamines.

2.3.6 2,4,5-T

Production of 2,4,5-T was carried out by an analogous process to that used for the manufacture of the 2,4-D. All components of the process were operated in the same manner, although at slightly different temperatures and pressures.

2.3.7 Esters of 2,4-D and 2,4,5-T

The esters of both 2,4-D and 2,4,5-T were formed by batch reaction with the appropriate alcohol in the presence of sulfuric acid which acted as a catalyst. The alcohol and by-product water were distilled throughout the reaction and the alcohol was held for later reuse in the process. On completion of the reaction, the last traces of water were removed under vacuum, and the ester was then transferred to storage to await sale.

2.3.8 Amines of 2,4-D and 2,4,5-T

The water-soluble dimethylamine (DMA) salts of 2,4-D or 2,4,5-T were formed by the reaction of a water solution of DMA with either wet or dry 2,4-D or 2,4,5-T. The resulting amine salt solutions were adjusted for concentration and then stored for later sale. A proprietary line of oil-soluble amines (the Dacamines) was prepared by the reaction of N-oleyl-1,3 propylenediamine with either dry, molten D or T acids or the flaked acids.

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2.4.2 Site Protection

An investigation conducted by NJDEP in May 1983 confirmed dioxin contamination as high as 51 ppm within the site boundaries. Following the discovery of dioxin on the site, Diamond Shamrock, at the direction of the NJDEP and EPA, took initial measures to control access to the property and reduce the possibility that dioxin-contaminated material could leave the property. The principal measures were:

- o A fence was installed around the property including the river front.
- o An around-the-clock security guard was placed at the only gate providing access to the property. The duty of the guard is to control entry onto the premises and restrict it to authorized personnel.
- o The entire site, excepting areas covered by buildings and equipment, was covered by a permeable geotextile fabric (Amoco No. 2002 polypropylene stabilization fabric). This fabric was weighted down by concrete blocks to prevent movement by wind.

Some portions of the fabric were disturbed during the sampling for the site evaluation field investigation. Repairs were made immediately to reestablish the protective integrity originally provided for the site.

2.5 SITE INVESTIGATION

Diamond Shamrock entered into an administrative consent order with the NJDEP on March 13, 1984. The order requires that Diamond Shamrock undertake certain actions to secure the site, prevent exposure to contaminants, and determine the vertical and horizontal extent of chemical contamination. The initial phase of the order requires the preparation of a work plan, the completion of a site evaluation, and a feasibility study of remedial alternatives.

Diamond Shamrock submitted a Work Plan for the initial phase to the NJDEP on April 18, 1984. As a result of NJDEP review comments, the Work

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TABLES

TABLE 2.2.1-1
RAW MATERIALS AND FINISHED PRODUCTS
DURING KOLKER, DIAMOND ALKALI,
AND DIAMOND SHAMROCK OWNERSHIP

Raw Materials

Acetic acid	Sulfuric acid
Acetic anhydride	Dimethylamine (40%)
*Acetaldehyde	Triethylamine
*Benzene	Chlorine
*Monochlorobenzene	2-Ethylhexanol
Tetrachlorobenzene	Butyl alcohol
*Chlorosulfonic acid	Isopropyl alcohol
Methanol	Butoxyethoxypropanol
*Oleum (20%)	*Nicotine
Phenol	Sodium Hydroxide

Finished Products

2,4,5-trichlorophenoxy acetic acid
2,4-dichlorophenoxy acetic acid
2,4,5-trichlorophenol
2,4,6-trichlorophenol
2,4-dichlorophenol
Monochloroacetic acid
*Hexachlorobenzene
*Dichlorodiphenyltrichloroethane
*p-chlorophenyl-p-chlorobenzene sulfonate (ovex)
*1,1,1-trichloroacetaldehyde
*Benzensulfonyl chloride
*p-chlorobenzenesulfonyl chloride
*p-chlorobenzenesulfonamide
*4,4'-dichlorodiphenylsulfone
*p-acetylamino benzene sulfonyl chloride
*p-methoxybenzene sulfonyl chloride
*1,2,4,5-tetrachlorobenzene
Amine salts of 2,4-D (dimethyl and triethyl amines)
Amine salts of 2,4,5-T (dimethyl and triethyl amines)
Esters of 2,4-3 (butyl, 2-ethylhexyl, isopropyl, butoxyethoxypropyl)
Esters of 2,4,5-T (butyl, 2-ethylhexyl, isopropyl, butoxyethoxypropyl)
Amine salts of N-oleyl-1,3-propylenediamine
*Nicotine sulfates
Muriatic Acid
*2,5-dichlorophenyl-p-chlorobenzene sulfonate

NOTE: The asterisk denotes raw materials and products not used or made after the explosion in February 1960

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TABLE 2.2.1-2
CHEMICALS PRODUCED BY KOLKER CHEMICAL WORKS

<u>Product</u>	<u>Estimated Total Production</u>
2,4-D	85 million pounds
2,4,5-T	25 million pounds
DOT	100 million pounds
HCB	10 million pounds
Ovex	10 million pounds
Lindane	Unknown
Low γ -BHC	<10 million pounds

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TABLE 2.2.2-1
MAJOR EVENTS DURING DIAMOND SHAMROCK AND
SUBSEQUENT OWNERSHIP

DATE	DESCRIPTION
3/51	Kolker Chemical Works, Inc., acquired by Diamond Alkali Co. Included in the transaction was the 1.8-acre parcel (the northeast portion of the site) on which the herbicides and DDT were being produced.
1/56	Easement obtained to construct a ten-inch sewer under the Central Railroad of New Jersey (now Conrail) tracks. A sewer connecting the plant to the PVSC sewer was subsequently installed. Following construction, most process wastes were diverted from the river to the PVSC treatment plant.
1957-58	Low γ - BHC production transferred to another location.
1958-59	DDT manufacture suspended following construction of a plant in Texas.
2/60	An explosion in the NaTCP unit destroyed the building in which it and several other products were made. Following this explosion, manufacture of all products other than the phenoxy herbicides and their intermediates was discontinued.
4/60	A 1.6-acre parcel (the southwest portion of the site) was leased from the Triplex Oil and Refining Co. (later Walter Ray Holding Co.) to permit expansion of the plant.
Mid 1961	Startup of new units to make NaTCP, DCP, and MCA.
1963	Rehabilitation of the 2,4-D acid process performed. Included was installation of improved ventilation in the area where 2,4-D and 2,4,5-T were made.
7/65	Startup of the melt, washing, and drying process to produce anhydrous, flaked 2,4-D. Capacity increased by 40 percent with this change.

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TABLE 2.2.2-1
(Continued)

DATE	DESCRIPTION
10/67	Construction of new 2,4-D unit and conversion of former 2,4-D unit to make 2,4,5-T completed. Both products were now made by the melt, washing, and drying process. Capacity increased by 54 percent for 2,4-D and 67 percent for 2,4,5-T. Purification of TCP also instituted as part of this expansion.
8/69	Production at the plant terminated by Diamond Shamrock.
9/69-12/69	All remaining raw materials and products were sold and removed from the site. Plant equipment was cleaned.
1/70-2/71	No activity. The plant was for sale.
3/16/71	Chemicaland Corp. purchased the 1.8-acre portion of the site and improvements from Diamond Shamrock, which also assigned rights to the lease of the 1.6 acres owned by Walter Ray Holding Co. to Chemicaland.
4/71-8/73	Cloray NJ Corp. (a Chemicaland affiliate) produced benzyl alcohol and some other specialties and performed custom manufacturing in the plant. Operations were suspended in August 1973.
9/73	Chemicaland contracted to make 2,4-D for Diamond Shamrock.
5/74-9/74	Limited quantities of 2,4-D were made during this period but none was delivered to Diamond Shamrock. Operations were suspended and the staff was laid off in September 1974.
2/75	Operations resumed by Chemicaland which was now making 2,4-D for Occidental Chemical Co.
11/22/76	Occidental Chemical assumed control of the management of the plant and continued to manage the plant until February 24, 1977, when control was returned to Chemicaland.
2/24/77	Chemicaland laid off all personnel and shut down the plant as things stood.
3/77-12/79	No information available on activities during this period.

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TABLE 2.2.2-1
(Continued)

DATE	DESCRIPTION
1980	The 1.8-acre parcel owned by Chemicaland was acquired by William Leckie (successor to Walter Ray Holding Co.), consolidating ownership of the whole site in his name.
3/81	The entire 3.4-acre site was sold to Marisol, Inc.
1981-82	Plant cleanup and clearing undertaken by Marisol.
1982-83	SCA Corporation leased part of plant and tank farm from Marisol.
5/83	Dioxin was found on the site. Cleanup action initiated by NJDEP and the EPA. Diamond Shamrock secured the site pending evaluation and remediation.

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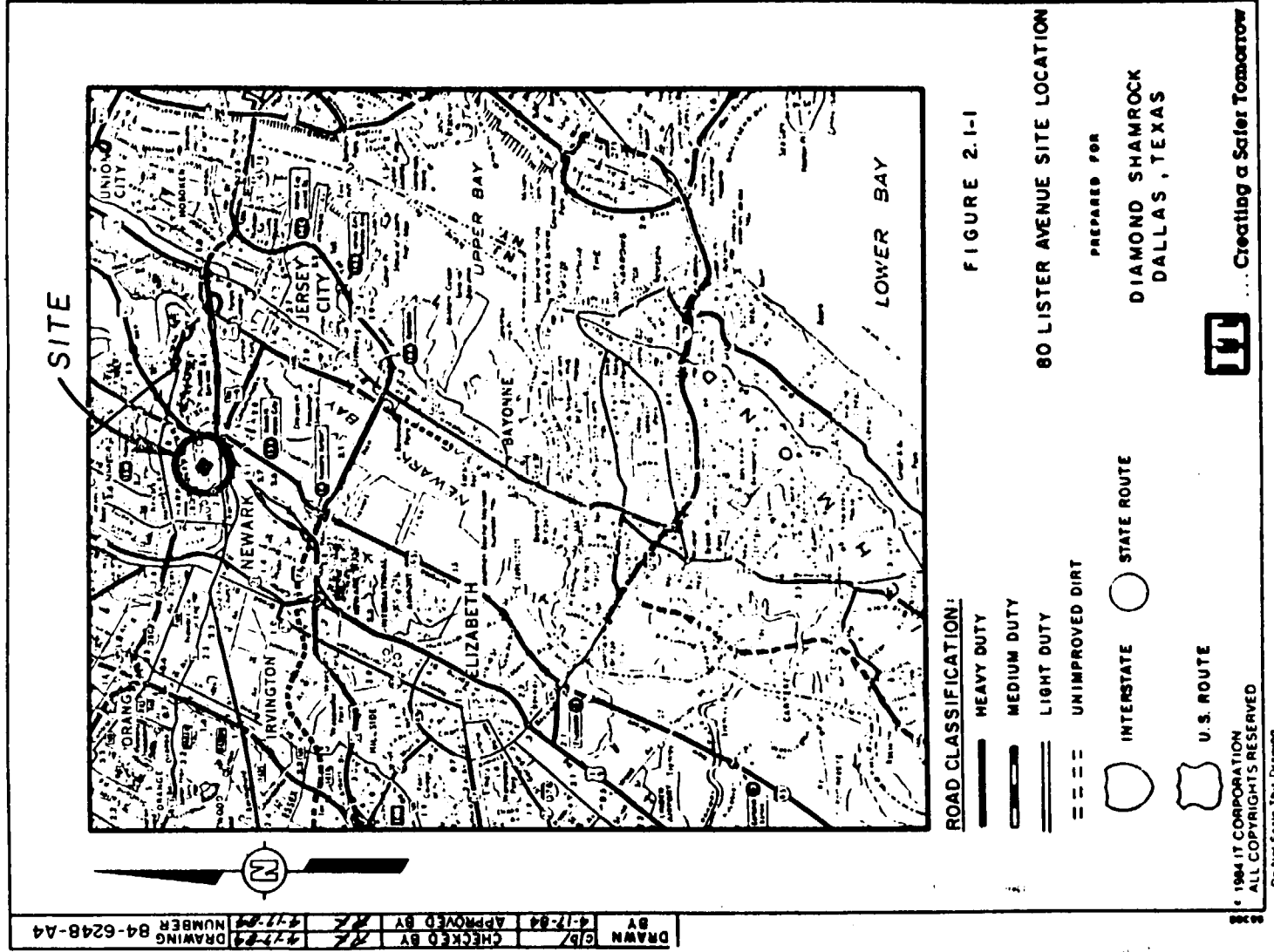
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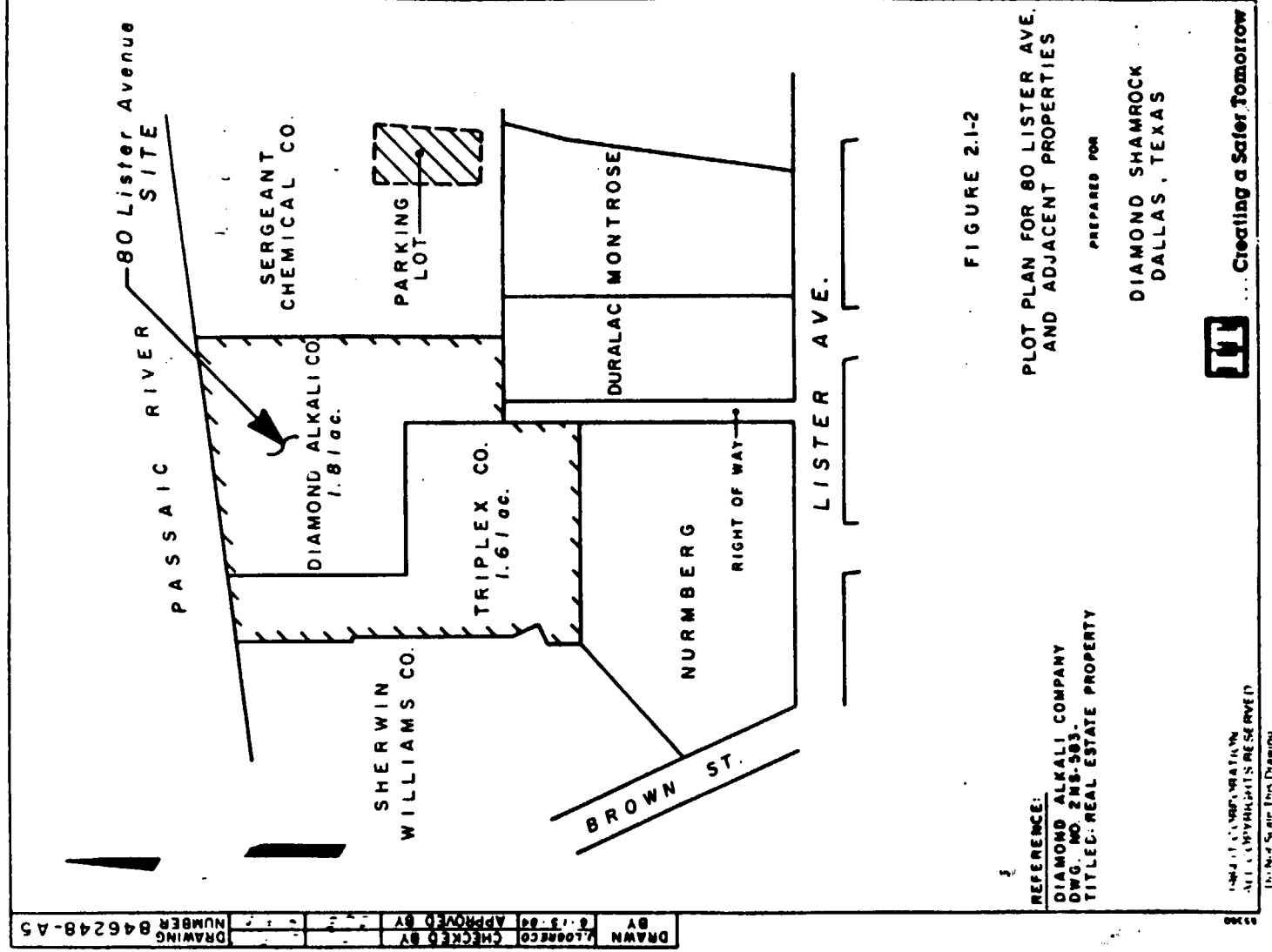
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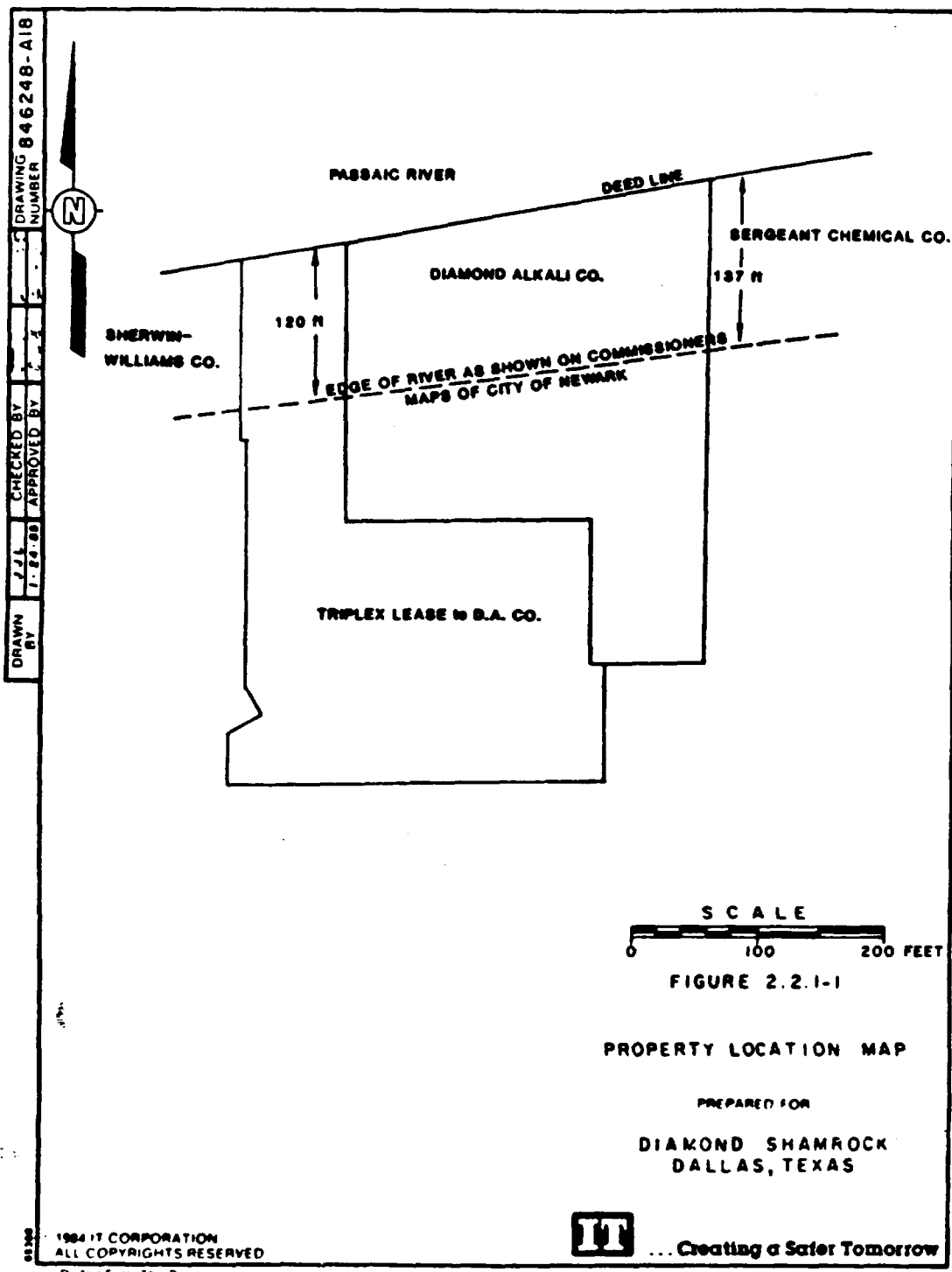
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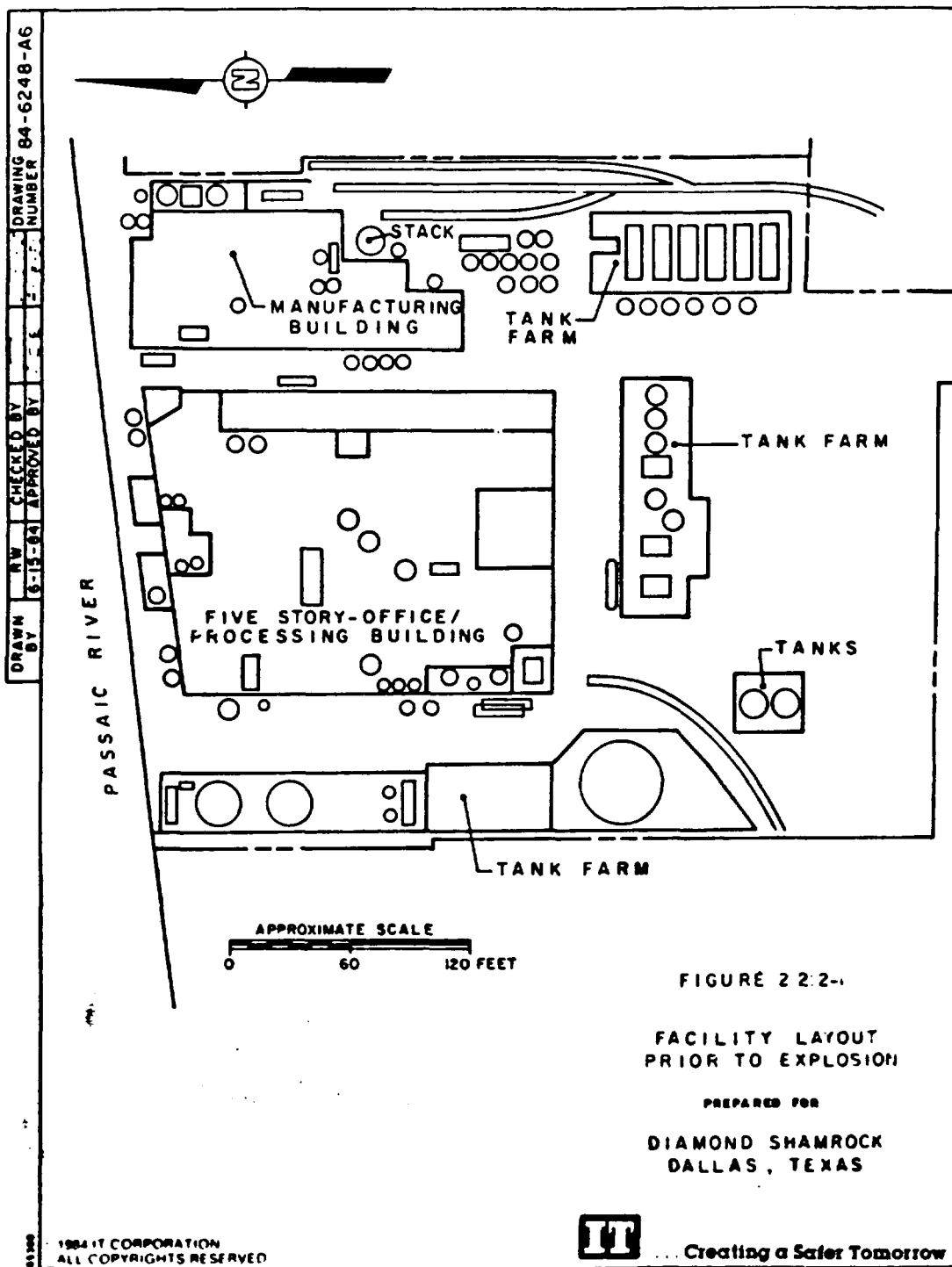


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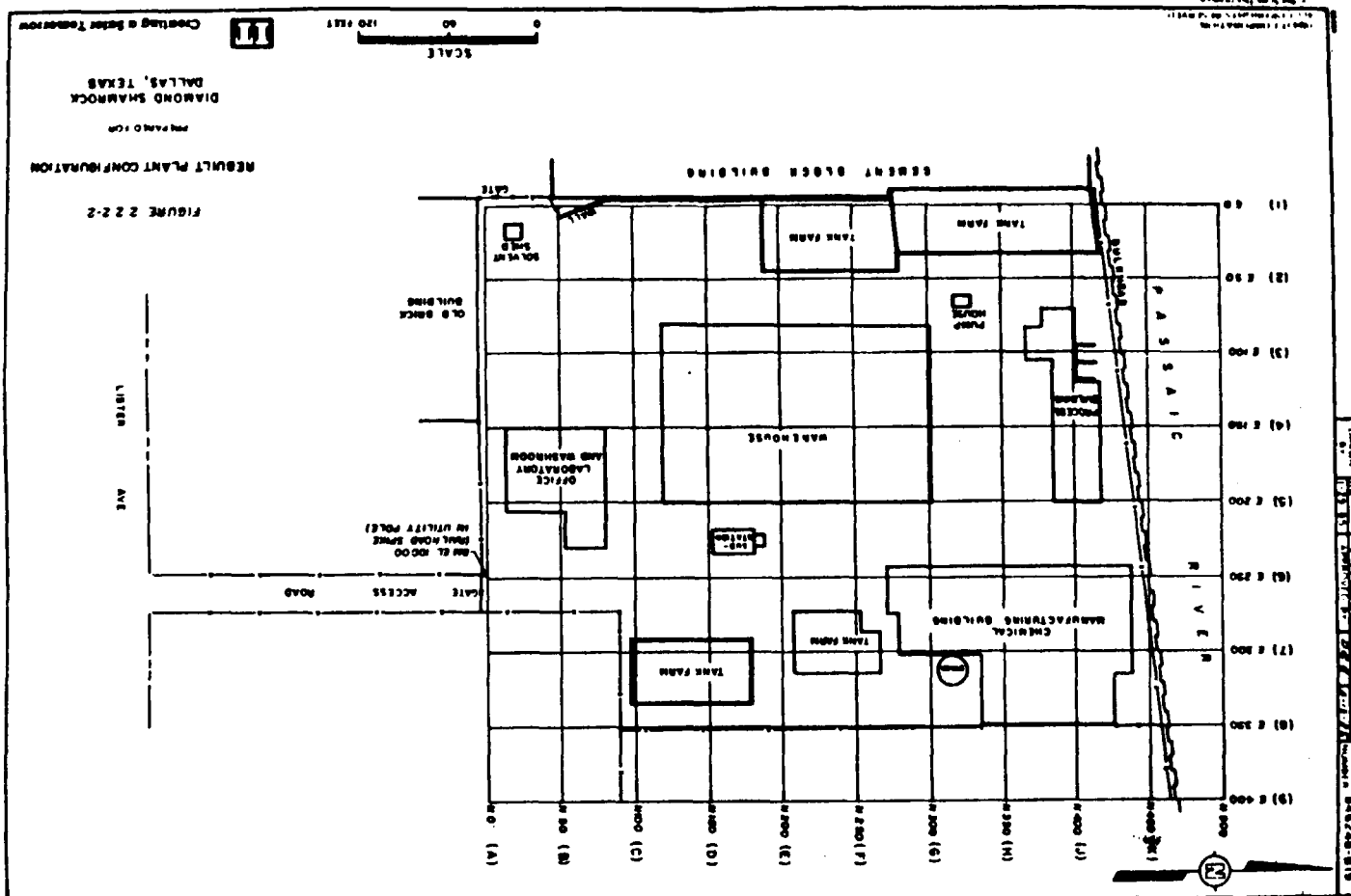


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3.0 REGIONAL SETTING

3.1 CLIMATE AND METEOROLOGY

3.1.1 General

Climate and meteorological conditions at the site have been characterized from 40 years of weather records available from the National Weather Service at Newark International Airport. The airport is located approximately three miles southwest of the site in a similar physiographic setting. The climatic data at the airport are, therefore, considered representative of conditions at the site.

The climate of the site is humid and is typified by moist, warm summers and moderately cold winters with winds of moderate velocity. The land around the site which was originally flat and marshy is now almost fully developed by industry and associated infrastructure.

3.1.2 Wind

Prevailing winds in the site area are from the southwest with only small seasonal variations in direction. Mean wind speeds are generally highest during the winter and spring months (10 to 12 miles per hour), while the lowest values (8 to 9 miles per hour) occur during the summer season. The predominant wind direction for winter months is west-northwest (13 percent of the time) while southwest (42 percent of the time) winds predominate during the summer. The highest wind speed recorded in the area was 82 miles per hour from the east in November 1950. During the 50-year period ending in 1965, 23 tornadoes were recorded in New Jersey resulting in two deaths and damage in the range of \$2.5 million (Dunlap, 1967).

The winds of the Newark area are greatly affected by two factors--the Atlantic Ocean and the topography of the surrounding area. To the northwest are ridges oriented roughly in a south-southwest to north-northeast direction. These ridges rise to an elevation of about 200

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(October) to 4.27 inches (August). Table 3.1.4-1 provides average precipitation data for Newark, including monthly and 24-hour maximums.

Precipitation patterns at the site are influenced by wind direction. Winds from either the north or the south bring most of the precipitation that falls in the area. Northerly winds consist of continental polar winds from Canada which collect moisture over the Great Lakes. This moisture is carried to northern New Jersey where it is deposited as rain or snow. When the wind is from the south, the warm, moist coastal winds bring moisture into the area.

3.1.5 Evaporation

Evaporation studies done in the area between 1956 and 1970 show that the average annual pan evaporation for Newark is 49.69 inches. Pan evaporation in Newark is highest in the month of July at 6.99 inches and lowest in the month of December at 1.64 inches. These evaporation values represent estimates occurring from a Class A pan (Farnsworth and Thompson, 1982).

Free water surface evaporation is the amount of water evaporated from a shallow lake, wet soil, or other moist natural surface. It is roughly 70 percent of the evaporation from a Class A pan for the same meteorological conditions. The annual free water surface evaporation for Newark is calculated to be approximately 35 inches.

3.2 GEOLOGY AND LANDFORMS

3.2.1 Physiography

The site is situated in the Piedmont Lowland section of the Piedmont Physiographic Province. This province is located between the Atlantic Coastal Plain and the Valley and Ridge Province (Figure 3.2.1-1).

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rocks were intruded into the sediments (Palisades sill, diabase dikes) and extruded onto the sediments (Watchung basalt flows) during the period of deposition.

The Newark Basin is bounded on the northwest by the Ramapo Border Fault. This fault is a normal fault which separates the Precambrian rocks of the highlands from the Triassic-Jurassic rocks of the lowlands. During the period of deposition and active tectonics, down-dropping of the basin continued along the fault. As a result, the rocks in the basin generally dip 5 to 25 degrees to the northwest. The oldest exposed formations are seen along the eastern edge of the basin and the youngest rocks are seen along the western edge at the fault. The total stratigraphic thickness of rock units deposited in the Newark Basin is calculated to be in excess of 30,000 feet.

The bedrock underlying the site is the Passaic Formation (Olsen, 1980), which is more commonly known as the pre-basalt portion of the Brunswick Formation (Kummel, 1940). This formation consists chiefly of soft red shale and sandstone.

Approximately 15,000 years ago, the area around the site was probably part of a delta (Figure 3.2.2-2) before being submerged beneath Lake Hackensack (Figure 3.2.2-3). The lake was fed by a retreating ice sheet to the north, dammed by a terminal moraine on the south, and confined between two erosional remnant ridges. About 10,000 years ago, the terminal moraine damming the lake was breached, resulting in the drainage of the entire body of water. The lake bed left behind developed into a flatland forest and a meadow now called the Mackensack Meadows.

The Mackensack Meadows lies between the First Watchung Mountain and the Palisades Ridge (Lovegreen, 1974) (Figure 3.2.2-1), and is a physiographic feature formed by sediment deposition in the bottom of the extinct glacial Lake Hackensack. The sediments consist of deposits of

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mean high water and mean lower water) is reported by NOAA as 5.1 feet. The spring range (average semidiurnal range occurring semimonthly as a result of the moon being New or Full) is reported by NOAA as 6.1 feet with the mean tide level (midway between mean low water and mean high water) at 2.5 feet.

The United States Geologic Survey (USGS) has developed equations using standard regression techniques for estimating mean annual flow of New Jersey streams. Because the Passaic River lies above the Fall Line (line of transition from the Atlantic Coastal Plain to the Piedmont Plateau), the USGS considers the river to be a noncoastal drainage basin. Flow in noncoastal drainages can be expressed as (USGS, 1984):

$$\begin{aligned} \text{mean annual flow (cfs)} &= 1.534 A \\ \text{where } A &= \text{drainage area in square miles} \\ \text{cfs} &= \text{cubic feet per second} \end{aligned}$$

Table 3.3.1-1 shows the mean annual flows for the Passaic River at the Little Falls gaging station and for the gaged tributary streams below Little Falls. The total drainage area of the Passaic River Basin is approximately 900 square miles which means that 55.5 square miles are not gaged in the lower basin. Using the USGS equation for noncoastal drainages, mean annual flow from the 55.5 square miles is calculated to be 85 cfs. Adding this flow to the total in Table 3.3.1-1 yields a mean annual flow at the mouth of the Passaic River of 1,392 cfs, or approximately 1,400 cfs.

The seven-day ten-year discharge (7-10) is the low flow that may be expected to occur on seven consecutive days on the average of once every ten years. A 7-10 value was estimated for the Passaic River at the project site from the Little Falls, New Jersey gaging station data and extrapolated by considering the additional drainage area between Little Falls and the site. Assuming the total drainage area is approximately 900 square miles, the 7-10 value has been estimated to be 68 cfs.

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Passaic River Dredging

A number of changes have been made to the river channel since the flood of 1903 that limit the severity of flooding that could occur from a similar storm. These factors include bulkheading and filling of the channel, dredging of the channel, and construction of dams on the river.

The Passaic River channel and the shipping channels in Newark Bay and the nearby Hackensack River are active commercial channels that are regularly maintained. The channels are dredged to permit navigation by barge and boat traffic. The dredging program is performed by the U.S. Army Corps of Engineers (COE). The COE (1975) has divided the navigable portion of the Passaic River into four separate sections of different project depths: the 10-foot, 16-foot, 20-foot, and 30-foot Projects (Figure 3.3.1-2).

The downstream limit of the 10-foot Project (channel dredged to a 10-foot depth) is approximately 8 miles upstream from the mouth of the Passaic. The 16-foot Project ends approximately 7 miles upstream from the mouth. The 20-foot Project extends to just downstream from the New Jersey Turnpike Bridge. The 30-foot Project includes the rest of the Passaic River to the confluence of the Passaic and Hackensack rivers.

The site is located on that section of the river designated as the 20-foot Project. In 1937 the Passaic River channel was dredged to a depth of 20 feet from the downstream limit of the 20-foot Project to the Jackson Street Bridge (Figure 3.3.1-2). Earlier, between 1921 and 1923, various portions of this segment of the channel were dredged to the project depth of 20 feet. Since 1937, however, no portion of the 20-foot Project downstream from the Jackson Street Bridge has been dredged (COE, 1984).

Because commercial traffic on the Passaic River above the 30-foot Project is declining, the COE has indicated that there may not be any reason

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and moves through and is stored in geologic formations. Geologic formations which can yield economically significant quantities of water to wells or springs are called aquifers. The regional aquifers in the vicinity of the site are the bedrock of the Brunswick Formation of Triassic age and the unconsolidated glacio-fluvial sands and gravel deposits of Pleistocene age.

The principal source of ground water in the area is the rocks of the Brunswick Formation. The shales and sandstones are generally capable of sustaining moderate to large yields to wells, but the Orange Mountain Basalt is capable of only small to moderate yields. The unconsolidated Pleistocene sand and gravel deposits, although capable of sustaining large yields, are of somewhat limited extent in the vicinity of the site.

Water in the rocks of the Brunswick Formation occurs under both unconfined and confined conditions. In the upland areas, the aquifer is generally unconfined. In the lowlands of the Hackensack Meadows, the aquifer is generally confined or semiconfined by glacio-lacustrine clay. Where the aquifer is confined by relatively impermeable layers, it is commonly under artesian pressure. The area around Newark has been subjected to heavy pumping, however, and the artesian pressure has been reduced. In part of Newark, extensive pumping has actually dewatered parts of the aquifer such that it no longer behaves as a confined aquifer (Nichols, 1968).

The ground water moves in the bedrock both vertically and horizontally from zones of secondary porosity through systems of interconnected joints and fractures. Most wells that are screened in this interval draw from more than one water-bearing zone, but the boundaries of the zones have not yet been accurately defined. Some wells penetrate from 400 to 600 feet below ground surface to reach these zones. The best producing wells, however are 300 to 400 feet deep (Nichols, 1968).

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Passaic River, the water levels by the year 1900 had been pumped to 40 to 130 feet below ground surface. Continued pumping in the 1900's has lowered the water level even further. This can be seen by comparing the contours in Figures 3.3.2-1 and 3.3.2-2. In 1879, evaluation of wells in the vicinity showed ground water levels from a few feet above to 25 feet below the ground surface. The heavy pumping has reversed the natural gradients in this vicinity and the dredging of the shipping channels in Newark Bay and the Passaic River has exacerbated the salt water intrusion problem by removing part of the barrier between the ground and surface waters (Nichols, 1968).

Ground water wells within a one-mile radius of the site were identified through permit records on file with the NJDEP. Table 3.3.2-1 presents the identified wells and their locations along with the relevant information obtained from the well permits.

3.4 FLORA AND FAUNA

The land in the vicinity of the site consists of tidal marsh and built-up land which is classified primarily as urban industrial. The terrestrial ecology of the natural environment is restricted to the tidal marsh, which has been modified by its proximity to the urban industrialized area. The industrialized area consists of a considerable number of buildings and an extensive amount of paved surface with very little exposed ground available to support flora or fauna.

Vegetation in the tidal marsh is primarily Phragmites australis (common reed), plus other wetland species such as Typha angustifolia (cattail) and Scirpus americana (bulrush). The terrestrial animals expected to be found in the immediate vicinity are likely to include various ground-feeding birds, eastern cottontail rabbits, and other small mammals such as the meadow vole. In the open marsh, muskrats are common and herptiles such as the garter snake and American toad are also present.

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- o Provide adequate supplies of industrial land uses which, in turn, will generate a broad range of job opportunities within the community.
- o Concentrate heavy industrial land use in the Meadowlands area; light industrial uses in both the Ironbound and the Triangle area (the area generally defined by Mulberry Street, Market Street, and McCarter Highway).
- o Allow for the controlled expansion of industrial development within areas where growth is now restricted due to incompatible abutting land uses.

The closest land area zoned for residential use in Newark is approximately one-quarter mile from the site.

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3.0 REFERENCES

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TABLES

TABLE 3.1.2-1
SEASONAL AND ANNUAL OCCURRENCE OF WIND DIRECTION

WIND DIRECTION	OCCURRENCE (Percentage of Time)				
	SUMMER	AUTUMN	WINTER	SPRING	ANNUAL
N	3.3	3.7	4	4.3	3.8
NNE	7	10	8	7	8.0
NE	5.7	6.3	6.7	6.7	6.4
ENE	3.7	3.7	3.3	5.7	4.1
E	3.7	3	3	5.3	3.8
ESE	4	3.7	2	6	3.9
SE	3.7	2.7	1	3	2.6
SSE	6.7	4	2	4.3	4.3
S	5.7	4	2.7	3.7	4.0
SSW	11.7	10	7	7	8.9
SW	12	11.7	9.7	8	10.4
WSW	8	8	9.3	7.7	8.3
W	5.7	7	7.7	5.7	6.5
WNW	6.7	7.3	13	9	9.0
W	5.7	6.7	11	8.3	7.9
WNW	5	6.3	9.3	7.7	7.1
	98.3	98.1	99.7	99.4	99.0

Reference: WCC, 1982

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TABLE 3.1.2-2
MONTHLY AVERAGE WIND SPEED
AND PREVAILING WIND DIRECTION (1)

MONTH	MEAN WIND SPEED (2) (mph)	PREVAILING WIND DIRECTION (3)
January	11.2	NE
February	11.5	NW
March	12.0	W
April	11.3	WNW
May	10.0	SW
June	9.3	SW
July	8.8	SW
August	8.6	SW
September	8.9	SW
October	9.3	SW
November	10.1	SW
December	10.8	SW
Annual	10.1	SW

(1) Recorded at Newark Airport.

(2) Length of record 35 years.

(3) Length of record 22 years.

Reference: WCC, 1982

TABLE 3.1.3-1
AVERAGE MEAN, MAXIMUM, AND MINIMUM TEMPERATURE
NEWARK, NEW JERSEY

(Data Period 1944 - 1983)

	MEAN (°F)	MAXIMUM (°F)	MINIMUM (°F)
January	31.4	38.5	24.3
February	32.5	40.1	24.8
March	40.9	49.0	32.7
April	51.4	60.7	42.0
May	62.1	71.7	52.4
June	71.2	80.6	61.8
July	76.5	85.6	67.3
August	74.8	83.7	65.9
September	67.7	76.7	58.6
October	56.8	66.0	47.5
November	46.1	53.9	38.2
December	35.1	42.2	28.0
Annual	53.9	62.4	45.3

Reference: NOAA, 1983

TABLE 3.1.4-1
PRECIPITATION DATA
NEWARK, NEW JERSEY

(Data Period- 1944 to 1983)

PRECIPITATION IN INCHES (Water Equivalent)

MONTH	NORMAL	MAXIMUM MONTHLY	YEAR	MAXIMUM IN 24 HRS.	YEAR
January	2.91	5.12	1964	1.78	1962
February	2.95	4.47	1956	2.45	1961
March	3.93	6.29	1954	2.58	1969
April	3.44	6.42	1958	2.01	1958
May	3.60	6.28	1968	4.11	1968
June	2.99	6.40	1975	2.31	1973
July	4.03	8.02	1975	3.40	1971
August	4.27	11.84	1955	7.84	1971
September	3.44	9.00	1975	5.27	1971
October	2.82	6.70	1955	3.04	1973
November	3.61	8.42	1972	3.78	1972
December	3.46	7.24	1973	2.14	1973
			Aug.		Aug.
Year	41.45	11.84	1955	7.84	1971

Reference: NOAA, 1983

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TABLE 3.3.1-1
DRAINAGE AREAS AND MEAN ANNUAL FLOWS
IN THE PASSAIC RIVER BASIN

STREAM AND STATION	DRAINAGE AREA (sq. mi.)	MEAN ANNUAL ⁽¹⁾ FLOW (cfs)
Passaic River at Little Falls	762	1,166
Saddle River at Lodi	54.6	98
Weasel Brook at Clifton	4.5	6
Third River at Passaic	11.8	22
Second River at Belleville	11.6	18
TOTAL	844.5	1,307

⁽¹⁾1980 U.S. Geological Survey Data

Reference: USGS, 1984

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Reference: New Jersey Department of Environmental Protection files

(1) TR_p = Brunswick Formation

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION (1)
Freeman Street Newark	P. Ballentine & Sons	1937	875	375	16	954	Industrial	TR _p
Lister Avenue	Kolker Chemical Works	12/11/51	802	600	12	1274	Industrial	TR _p
Lister Avenue	Kolker Chemical Works	4/27/49	359	300	10	984	Industrial	TR _p
171 Blanchard St.	Eureka Construc- tion Co.	1/23/59	500	75	8	904	Industrial	TR _p
Newark	Cajane Corp. of America	5/16/47	856	778	16	71	Industrial	TR _p
1215 Harrison Ave. Kearny	Theobald In- dustries	1973	584	350	12	85	Industrial	TR _p
Alh & Passaic River Harrison	PSE&C	1932	804	406	12	218	Industrial	TR _p

PERMITTED WELLS WITHIN ONE MILE OF 80 LISTER AVENUE

TABLE 3.3.2-1

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Reference: New Jersey Department of Environmental Protection files

(1) TR_p = Brunswick Formation

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION (1)
1180 Raymond Blvd. Newark	Newark Center Corp.	1955	700	89	10	147	Industrial	TR _p
Harrison	D. L. and W&R	1952	225	18	6	112	Industrial	TR _p
196 Blanchard St. Newark	International Metallurgical	1980	300	150	6	72	Domestic	TR _p
12 Lister Avenue Newark	ACMC Refinery Company	1960	500	150	10	140	Industrial	TR _p
17 Blanchard St. Newark	Newark Box Board Company	1981	400	105	10	72	Industrial	TR _p
117 Blanchard Ave. Newark	Fairmount Chemical	1968	250	200	8	-	Industrial	-
117 Blanchard Ave. Newark	Fairmount Chemical	1965	300	300	8	66	Industrial	TR _p

TABLE 3.3.2-1
(Continued)

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Reference: New Jersey Department of Environmental Protection files

(1) TR_p = Brunswick Formation

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION (1)
70 Blanchard St. Newark	Newark Paraffin Co.	1968	603	100	8	65	Industrial	TR _p
60 Blanchard St. Newark	Arden Chemical	1962	400	90	8	902	Industrial	TR _p
Brill Street Newark	Standard Bitulithic Co.	1964	406	360	10	73	Industrial	TR _p
325 Raymond Blvd. Newark	Hildeman In- dustries	1981	400	38	6	-	Industrial	TR _p
45 Manufacturers Place, Newark	Bonson Metal Corporation	1981	400	150	8	80	Industrial	TR _p
45 Manufacturers Place, Newark	Bonson Metal Corporation	1965	300	220	8	75	Industrial	TR _p
45 Manufacturers Place, Newark	Bonson Metals Corporation	1973	300	150	8	82	Industrial	TR _p

TABLE 3.3.2-1
(continued)

TABLE 3.3.2-1
(Continued)

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION ⁽¹⁾
183 Foundry St. Newark	Arkansas Company	1965	400	65	8	-	Industrial	TR _b
50 Paris St. Newark	Federal Pacific Elec. Prod.	1955	500	250	10	120	Industrial	TR _b
18 Avenue "L" Newark	John Englehorn & Sons	1952	500	450	10	-	Industrial	TR _b
84 Foundry St. Newark	Pfuff & Kendall	1965	200	100	6	-	Industrial	TR _b
Lincoln Hwy. Kearny	Coca-Cola Co.	1981	650	20	6	95	Domestic	TR _b
354 Doremus Ave. Newark	Celanese Chemical Company	1981	700	100	-	-	Industrial	TR _b
260 Schuyler Ave. Kearny	Standard Plastics	1974	250	150	6	105	Industrial	TR _b

(1) TR_b = Brunswick Formation

Reference: New Jersey Department of Environmental Protection files

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TABLE 3.3.2-1
(Continued)

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION ⁽¹⁾
574 E. Ferry St. Newark	Technical Plastic Extruders, Inc.	1981	300	422	8	85	Industrial	TR _b
244 Dukas St. Kearny	WilPet Tool Manu- facturing Co.	1961	700	520	10	-	Industrial	TR _b
Bergen Ave. Kearny	Nick Versaleno	1959	235	150	6	145	Industrial	TR _b
Harrison	Reynolds Metal Co.	1941	467	350	8	78	Industrial	TR _b
100 Schuyler Ave. Kearny	C&A Exxon	1982	25.5	-	4	-	Observation Well	Sand
100 Schuyler Ave. Kearny	C&A Exxon	1982	22	-	4	-	Observation Well	Sand
100 Schuyler Ave. Kearny	C&A Exxon	1982	23	-	4	-	Observation Well	Sand

(1) TR_b = Brunswick Formation

Reference: New Jersey Department of Environmental Protection files

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Reference: New Jersey Department of Environmental Protection files

(1) TR - Brunswick Formation

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION (1)
Harrison & 7th St. Harrison	Town Park Exon	1982	33	-	4	-	Observation Well	Sand
Harrison & 7th St. Harrison	Town Park Exon	1982	32	-	4	-	Observation Well	Sand
Harrison & 7th St. Harrison	Town Park Exon	1982	32	-	4	-	Observation Well	Sand
Harrison Ave. & Ann St. Harrison Baking		1981	30	-	4	-	Monitoring Well	Sand
Harrison Ave. & Ann St. Harrison Baking		1981	30	-	4	-	Monitoring Well	Sand
Harrison Ave. & Ann St. Harrison Baking		1981	30	-	4	-	Monitoring Well	Sand
Harrison Ave. & Ann St. Harrison Baking		1981	30	-	4	-	Monitoring Well	Sand
Harrison Ave. & Ann St. Harrison Baking		1981	30	-	4	-	Monitoring Well	Sand
Pennsylvania Ave. Kearny	Honsence	1983	10	-	4	-	Monitoring Well	Sand

TABLE 3.3.2-1 (Continued)

TABLE 3.3.2-1
(Continued)

REPORTED ADDRESS	OWNER	DATE DRILLED	TOTAL DEPTH BELOW SURFACE ELEVATION (ft)	YIELD (gpm)	DIAMETER (in)	DEPTH BELOW SURFACE ELEVATION TO BEDROCK (ft)	USE	FORMATION ⁽¹⁾
Pennsylvania Ave. Kearny	Monsanto	1983	10.5	-	4	-	Monitoring Well	Sand
Pennsylvania Ave. Kearny	Monsanto	1983	9.5	-	4	-	Monitoring Well	Sand
Pennsylvania Ave. Kearny	Monsanto	1983	10	-	4	-	Monitoring Well	Sand
Pennsylvania Ave. Kearny	Monsanto	1983	10	-	4	-	Monitoring Well	Sand
300 ft. W. of Jacobus 1/4 mi. W. of Pulaski Skyway	Syncon Corp.	1976	403	-	10/8	-	-	-

(1) TR₁ = Brunswick Formation

Reference: New Jersey Department of Environmental Protection files

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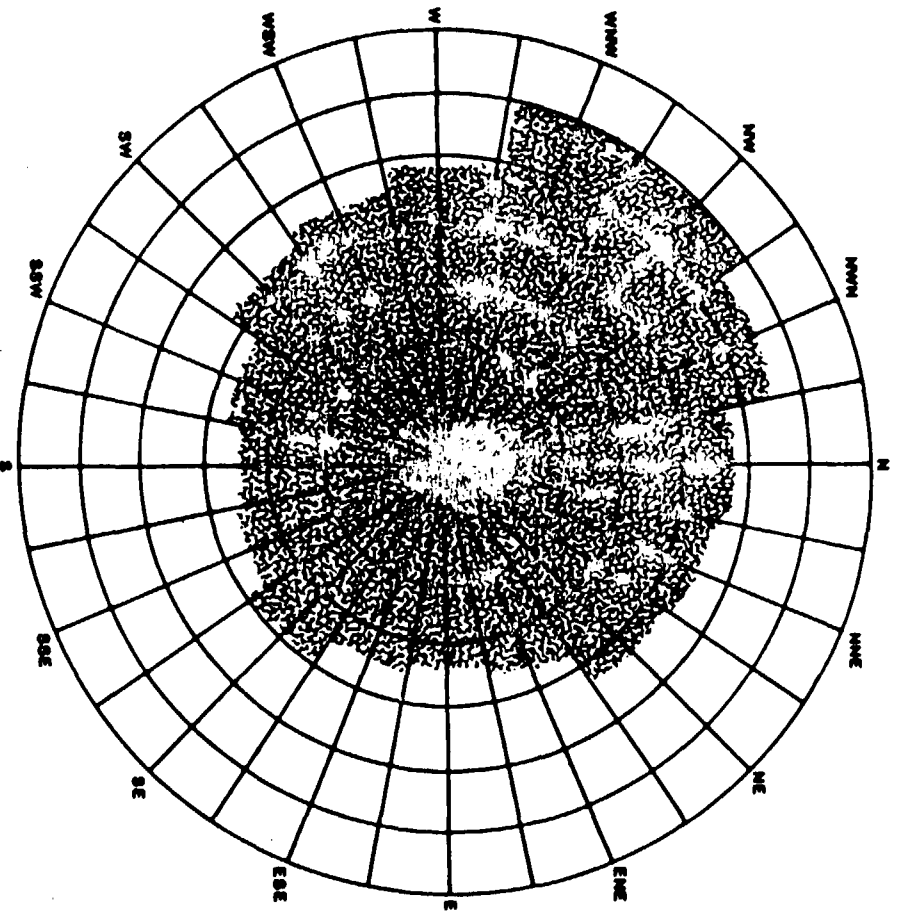
FIGURES

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1974

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	1-24-83	APPROVED BY			



REFERENCE: NOAA, 1978

FIGURE 3.1.2-2
AVERAGE WIND SPEED
FOR WIND DIRECTION
1965 - 1974
PREPARED FOR
DIAMOND SHAMROCK
DALLAS, TEXAS



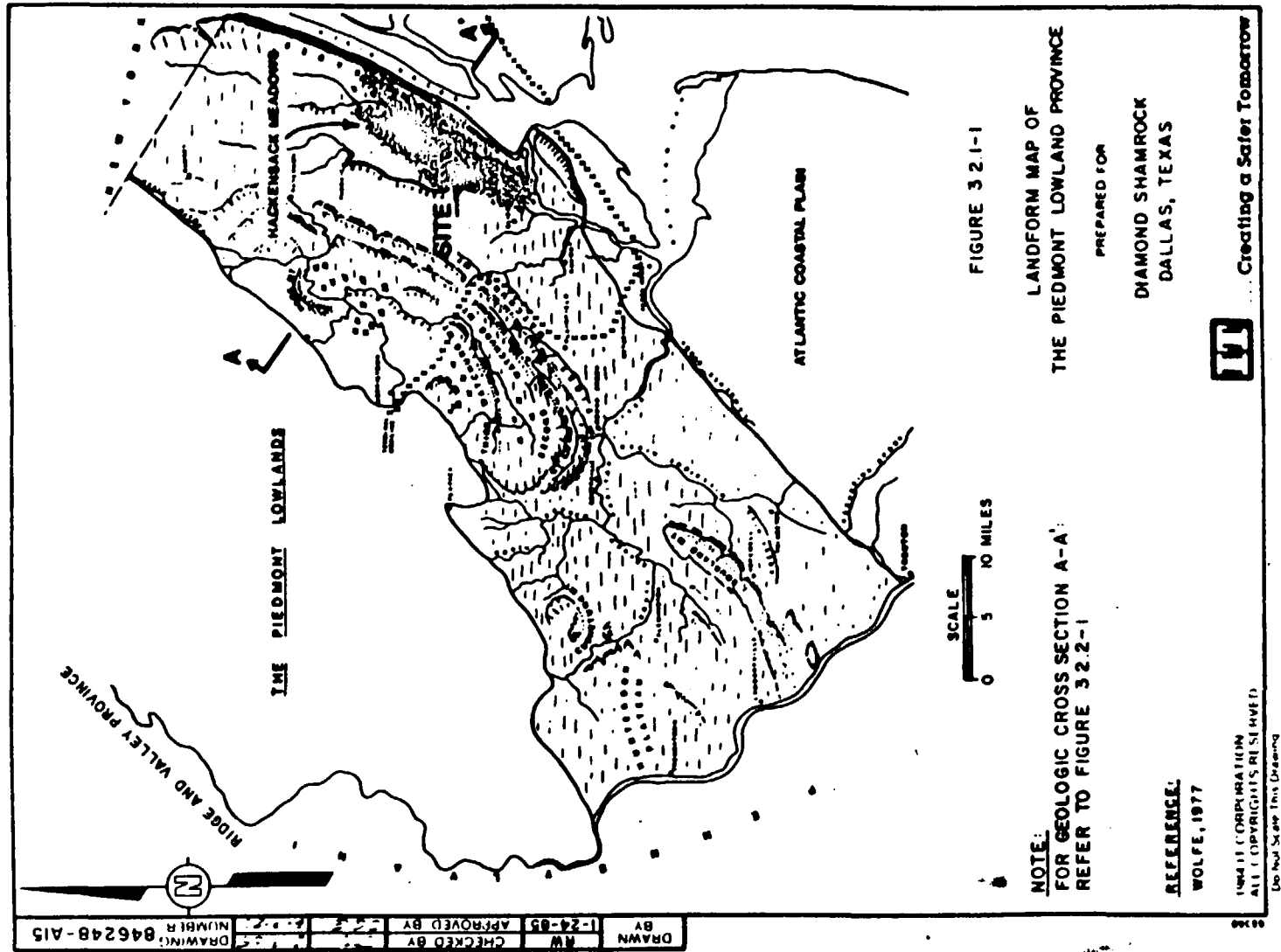
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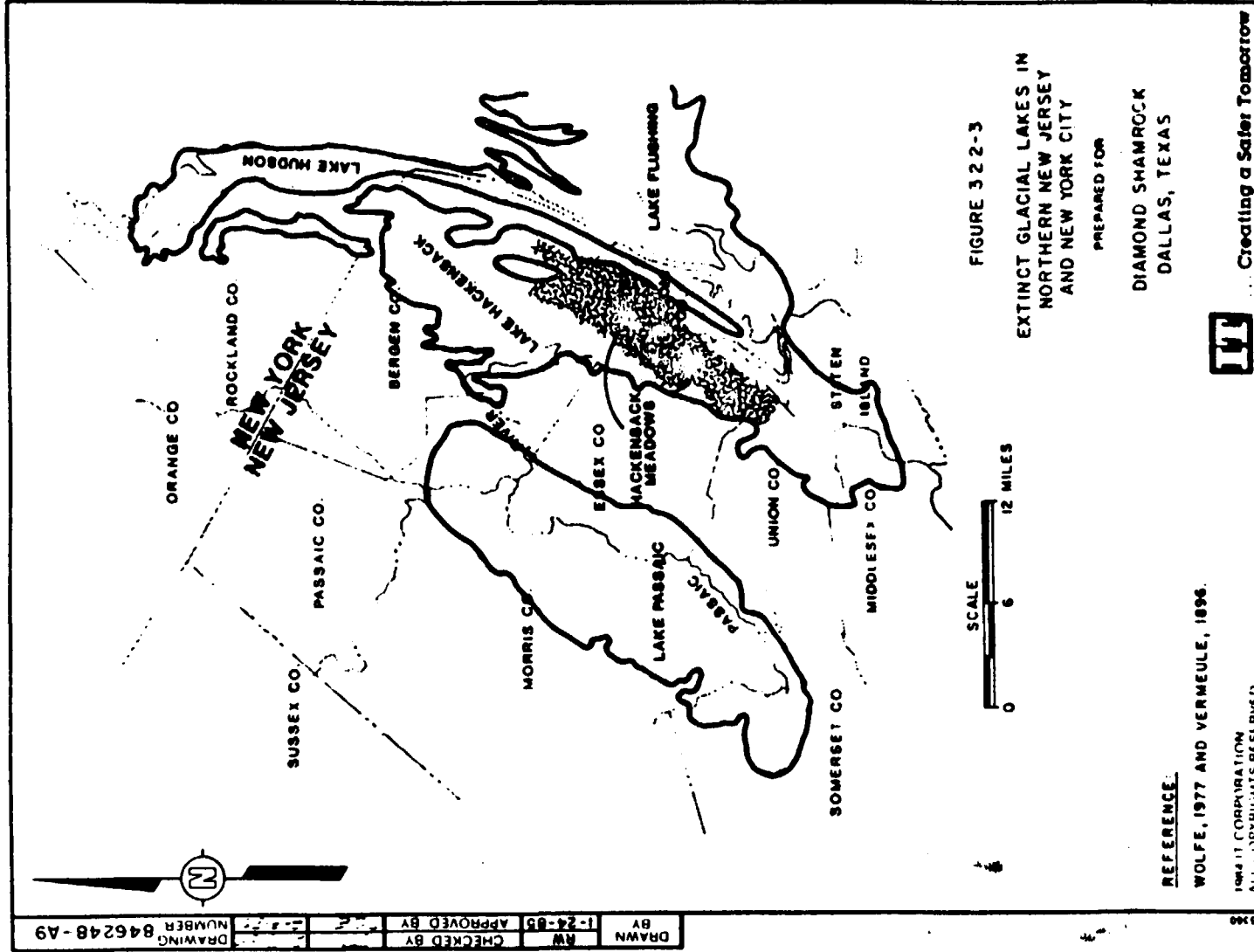
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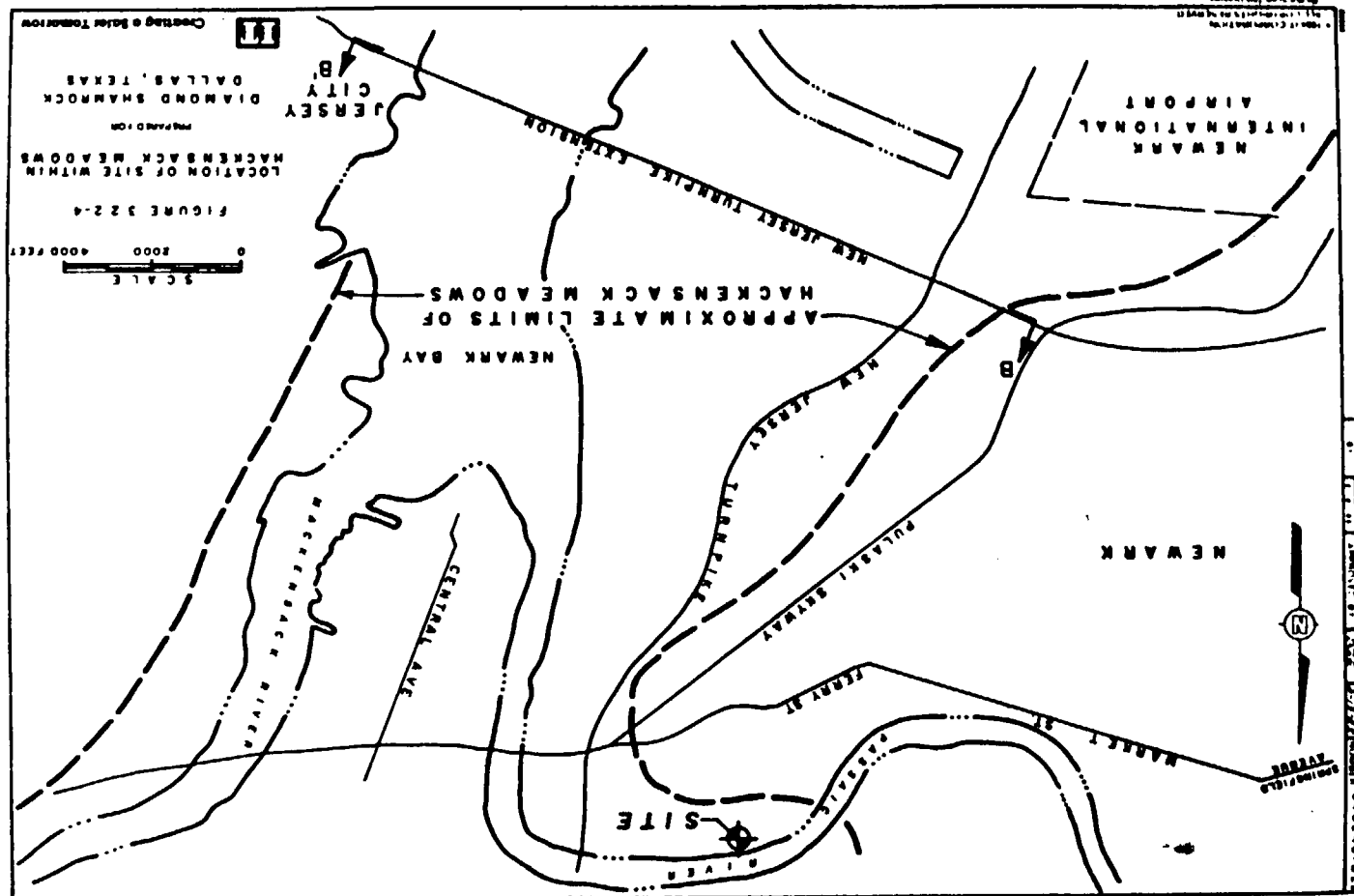
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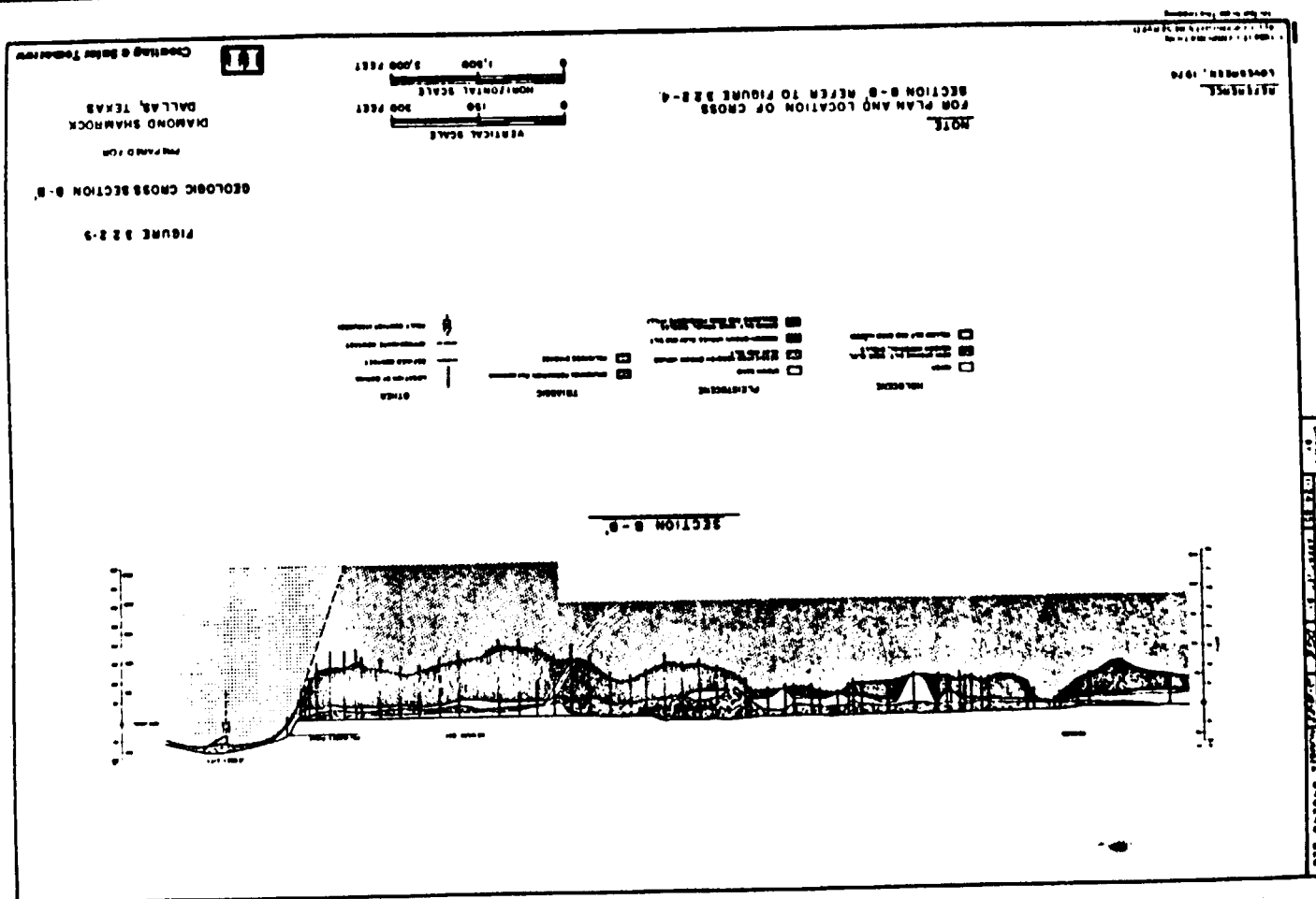
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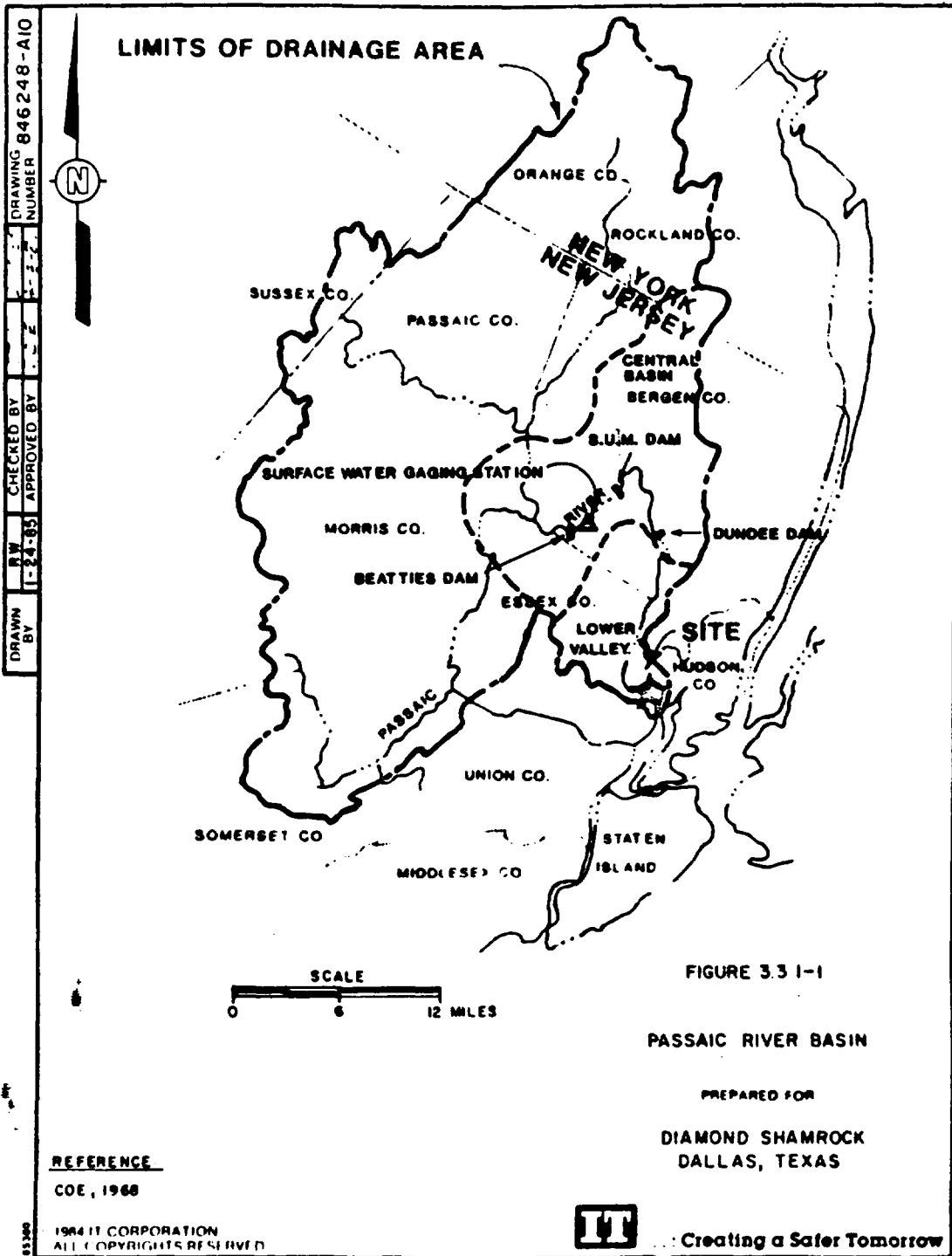
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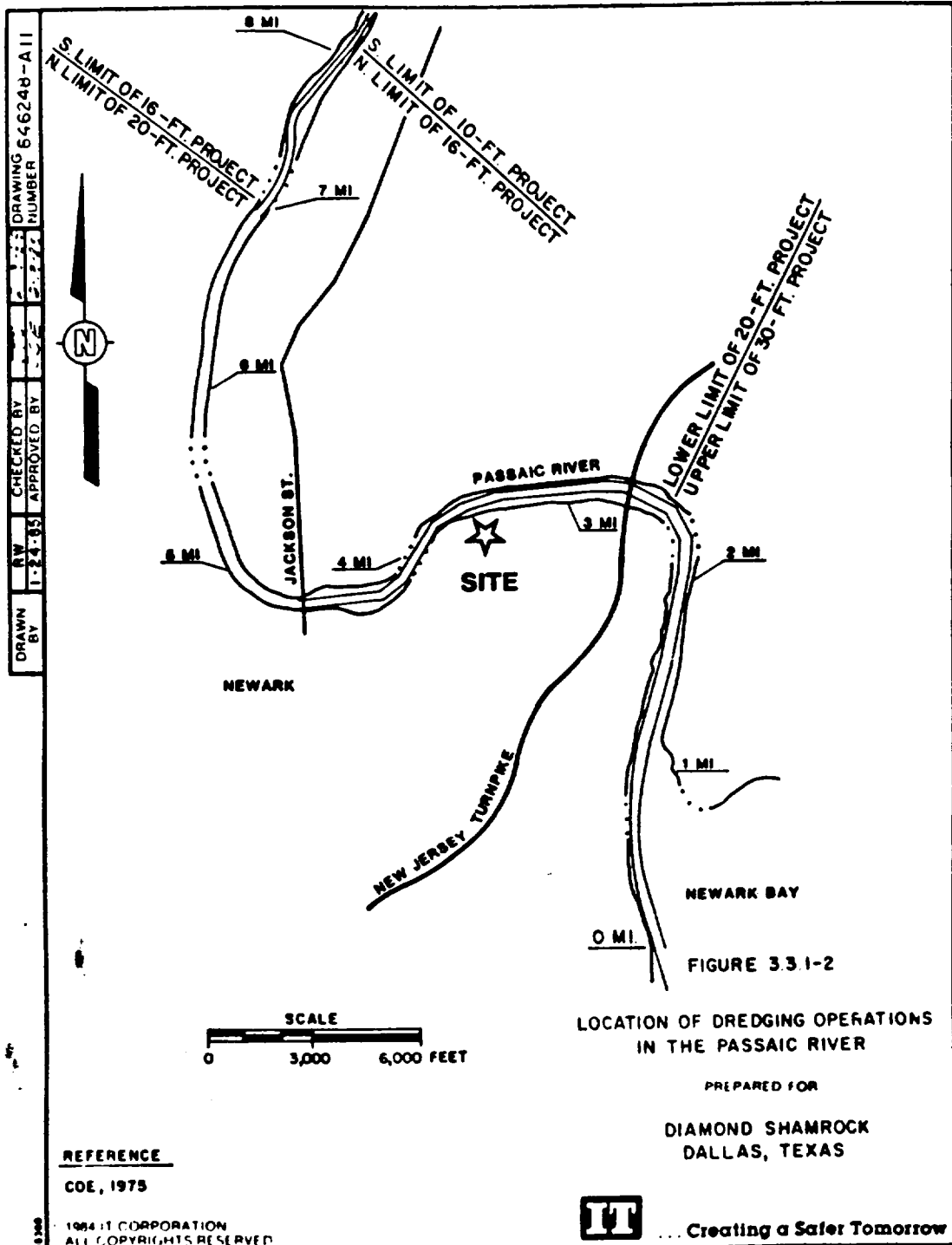


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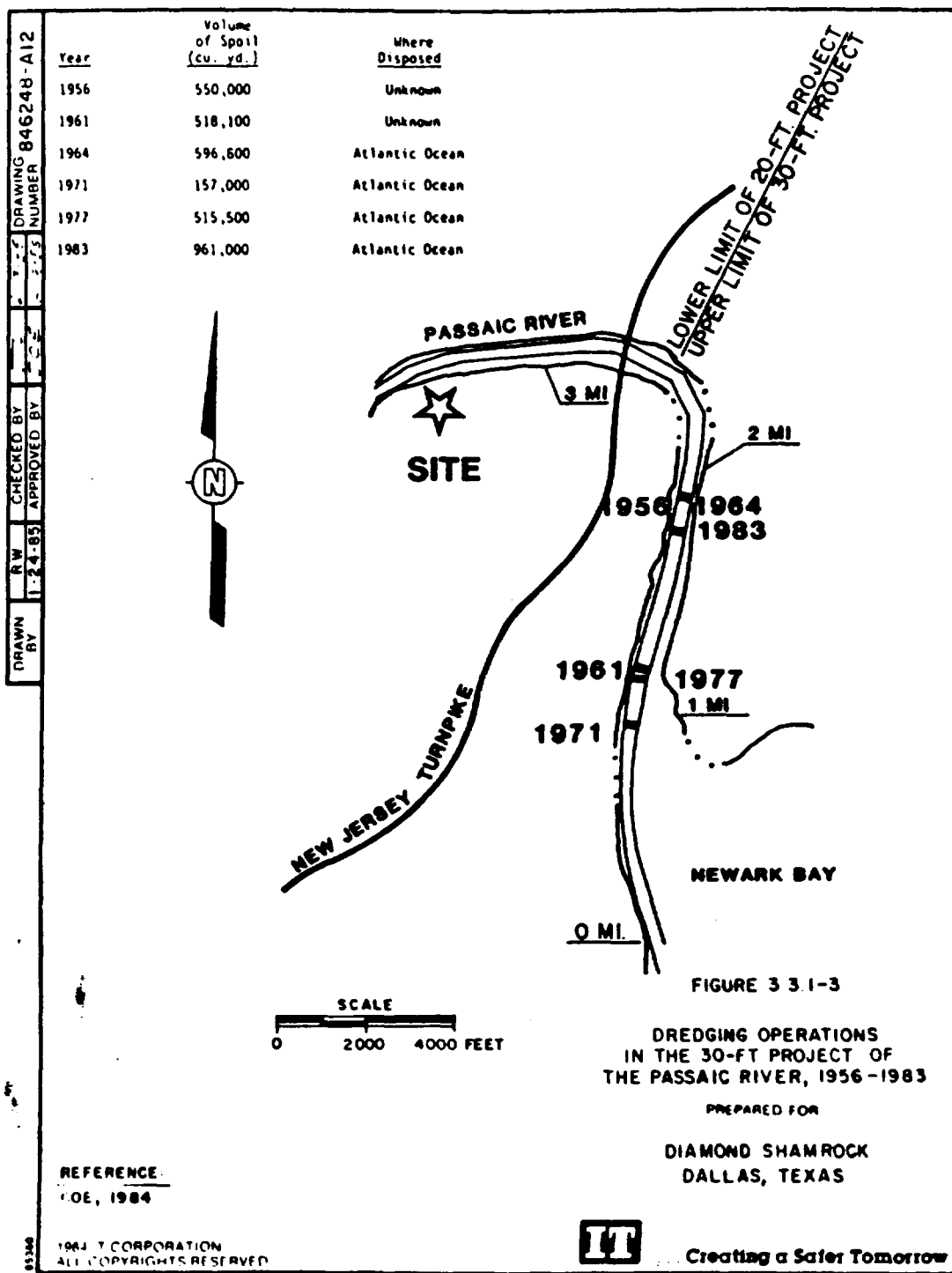
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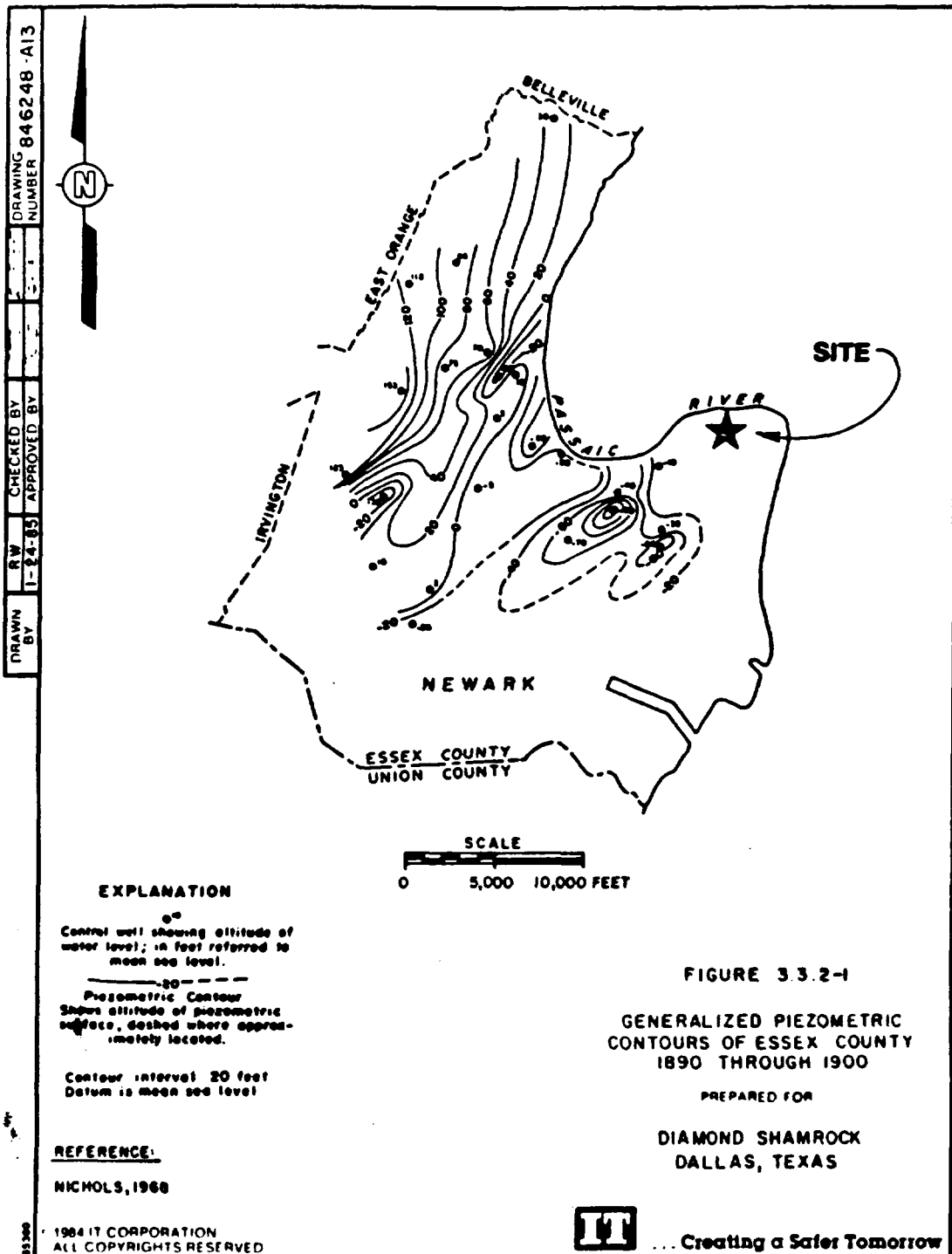
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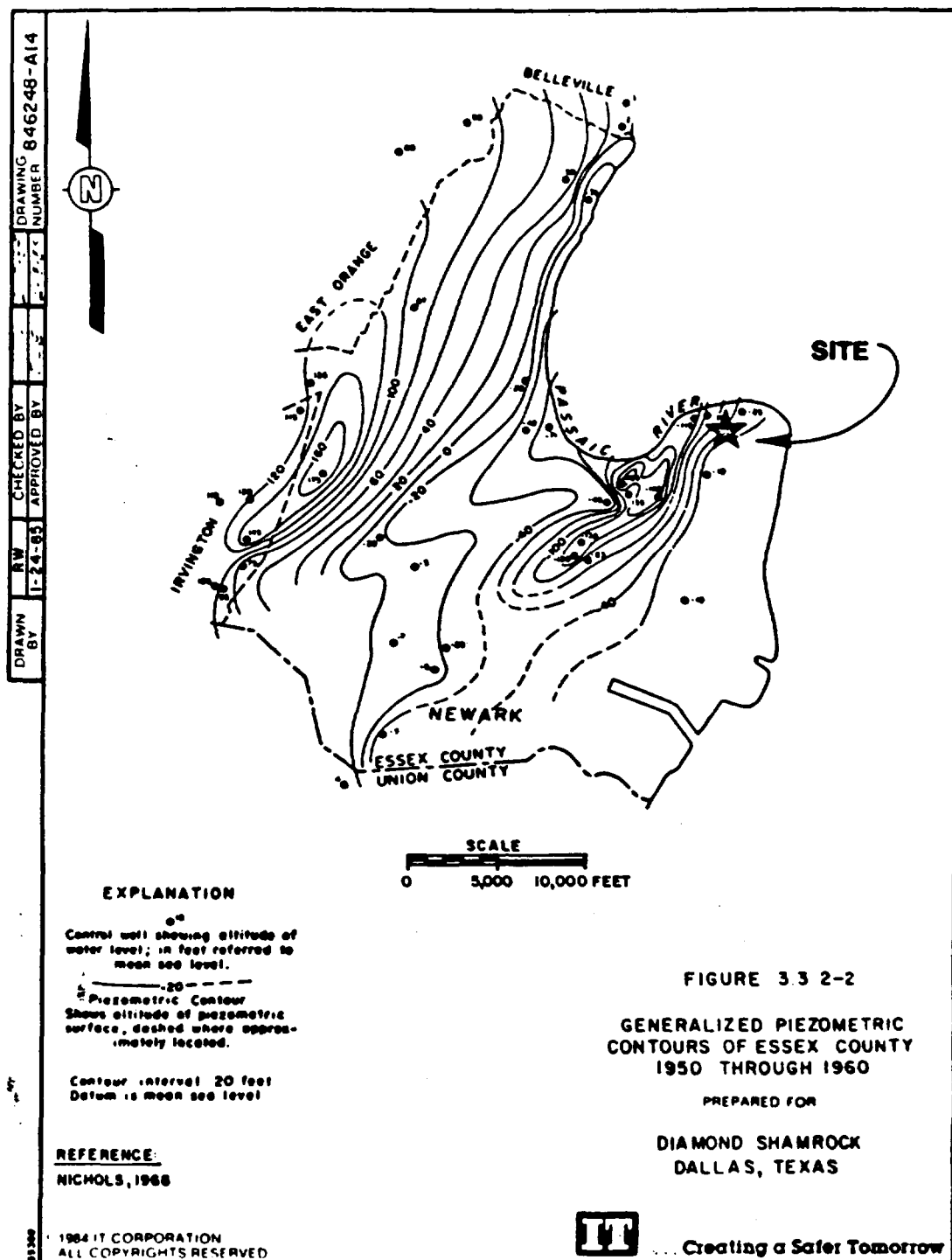
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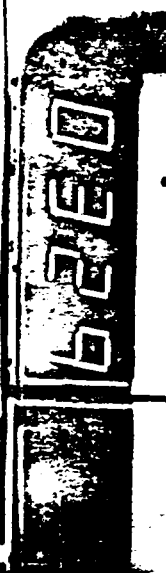
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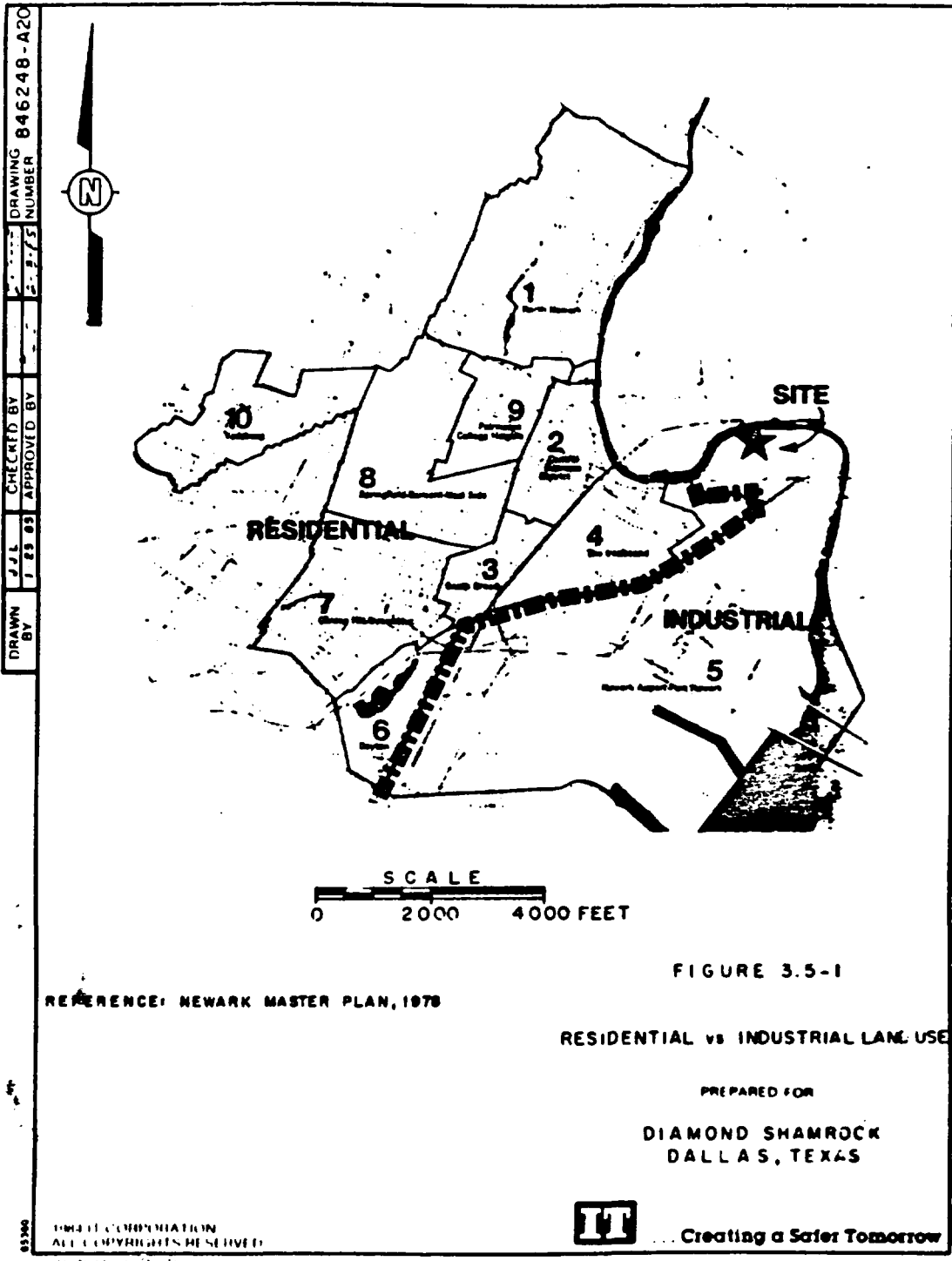


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4.0 SITE INVESTIGATION

A comprehensive field investigation and sampling program was developed for the site evaluation. Major considerations in the development of the program included: sampling and analytical testing methodologies; sample collection, handling, documentation and transportation; initiation of a health and safety program to protect project personnel and the general public; and quality assurance/quality control (QA/QC) protocol.

The selection of specific sample locations for soils, buildings and structures was concentrated at those points on the site considered to have the highest potential for the presence of dioxin and/or other contaminants. This biased sampling approach, based on plant historical data, was considered to provide the best data base available for a realistic overall evaluation of the extent of site contamination. A plan of the site showing major buildings and facilities and other physical features is provided in Figure 4.0-1.

A variety of sampling activities was performed to characterize the levels of chemical contamination at the site and to meet required health and safety and QA/QC requirements. These included:

- o Ambient air samples
- o Industrial hygiene samples
- o Chip, wipe, and bulk samples from existing buildings, tanks, piping, equipment, and sewers
- o Samples of soil
- o Samples of ground water
- o Samples of Passaic River water and sediments
- o Samples of background soil
- o Samples of on-site drums

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sequential numbers were assigned to all drums, regardless of their location.

The four-digit sample location numbers were assigned by the Sampling Coordinator before sampling personnel were sent into an assigned area, and this information was recorded in a field log notebook.

Additional information detailing sample identification is provided in Appendix A.

Key project personnel and their major duties and responsibilities during the site investigation included:

- o Field Operations Manager - Overall supervision of daily field activities, client and regulatory agency liaison, and responsibility for adherence to programs and schedules.
- o Site Manager - Administration of site personnel and equipment, supervision of equipment, and materials purchases and maintenance.
- o Sampling Coordinator - Supervision of sample handling staff and responsibility for maintaining proper sample handling techniques, sample shipment, and documentation of sample records.
- o Task Supervisors - Supervision of field personnel assigned to specific tasks (i.e., chip samples, drums, near-surface soil samples, etc.) and responsibility for field documentation of these tasks.
- o Site Health and Safety Coordinator - Responsibility for the administration of on-site health and safety programs, personnel monitoring, and decontamination procedures.
- o QA Program Manager - Responsibility for supervision and documentation of QA/QC program.

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- o A procedure to comply with the New Jersey Worker and Community Right to Know Act.
- o An Emergency Action Plan which provided detailed procedures in case of an accident or illness on site.

At least one industrial hygienist was on site at all times to coordinate the health and safety program.

4.1.2 Sample Handling and Documentation

A project-specific Standard Operating Procedure (SOP) was developed for all sample handling and documentation. This SOP is provided in Appendix A, and describes in detail field sample collection, the unique identification of samples, the shipping of samples to laboratories for analysis, the documentation associated with all these activities, and the reporting and permanent filing of the analytical laboratory results. Attachments 1 and 2 to Appendix A provide detailed information on sample identification codes and procedures for field log notebook entries, respectively.

A brief summary of sample handling and documentation procedures is provided in the following paragraphs.

Each day's sample collection was planned in advance by the Field Operations Manager. The Sampling Coordinator then assigned an alphanumeric code designation to each sample, and entered them into the Master Sample Collection and Shipping Log. The code designation indicated: sample origin (relative to the site grid), sample type, depth (where appropriate), and laboratory destination(s). Each sample collected was logged into a field log notebook by the person obtaining the sample. A label with the alphanumeric code was attached to the page describing the sample collection.

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Some adjustments to the QA Plan were necessary at the beginning of the project due to changes in the scope or objectives of the program, logistical problems, or requests by the NJDEP. These adjustments included redefinition of the collection schedule for sampling equipment rinsates, field blanks, and trip blanks; reassessment of preservation needs and shipment requirements; and revision of the sample custody procedures. The revised QA/QC procedures instituted for these areas are described in the following subsections. In addition, the details of the QC procedures followed during preparation of sample containers are provided. Finally, a description of the requirements for QC check samples at the participating laboratories is presented, and the QC checks initiated at the program level are described.

4.1.3.1 Field and Trip Blank Requirements

For this project, a field blank was defined as laboratory-pure water poured over a piece of sampling equipment and caught in an empty sample container on site. A field blank was obtained by a sampling team member immediately prior to the start of each day's collections. The field blank is identical to the "equipment rinsates" described in the original QA Plan. The water and containers for field blanks were provided daily to the sampling teams by the Sampling Coordinator, who received them from the Middlebrook Pike laboratory.

Trip blanks consisted of laboratory-pure water provided to the sampling teams in sealed sample containers; the containers were filled at the Middlebrook Pike Laboratory, labeled by the on-site Sampling Coordinator, and issued daily to the sampling teams. They were handled in exactly the same manner as the field samples, but the containers were not opened on site for any reason.

The frequency requirement requested by the NJDEP for field and trip blanks, was one field blank and one trip blank per day. Blanks associated with soil and river sediment samples were routinely analyzed for

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returned to the Sampling Coordinator with the samples. As the field sampling personnel released the samples to the sample handling personnel, they signed the Chain-of-Custody records, marking the first custody transfer.

4.1.3.4 Sample Container Preparation

All sample containers used at the site were supplied by the Middlebrook Pike laboratory. The containers were cleaned prior to shipment to the site according to the procedure described in the Work Plan. One container from each set of 40 was rinsed with laboratory-pure water, and the rinsate was analyzed for the same parameters as the sample to be collected in that type of container. Lot designations indicating the check analysis performed were assigned to each set of 40 containers and these were written on the boxes, indicating the check analysis performed:

DX	--	Dioxin
AA	--	Metals
PP	--	All PP parameters (used for soil containers)
EO	--	Extractable Organics
PO	--	Purgeable Organics
CN	--	Cyanide
OH	--	Phenols.

The last two characters of the lot designation indicate the lot number. As containers were issued to sampling teams on site, the lot designations were recorded in the Master Sample Collection and Shipping Log. Records including preparation dates, analytical data, and final results verifying container quality were filed with the project data.

4.1.3.5 Laboratory Quality Control Checks

Samples collected from the site were shipped directly to one or all of the three participating IT Analytical Services laboratories for analysis. EPA Contract Laboratory Program (CLP) methods were used for analysis of dioxin and the organic priority pollutants, as described in the Work Plan, with stated modifications in the case of the dioxin

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4.1.3.6.1 Non-VOA Field Blanks

Routine field blank requirements included analysis of volatile organics for blanks associated with soil and sediment samples only. Between October 9 and October 23, 1984, 10 additional field blanks were randomly assigned in conjunction with the usual VOA blanks for analysis of metals, cyanides, phenols, semivolatiles, and dioxin (two blanks for each parameter).

4.1.3.6.2 Blanks

A set of three unused gauze wipes was prepared at the Directors Drive laboratory as blank samples. They were assigned label numbers by the QA Program Manager and inserted into the on-site schedule for shipment with routine site samples to the laboratory for analysis of dioxin. Similarly, a set of three soil samples (Clarksburg soil, clean with respect to dioxin, originally obtained from Mason & Hanger-Silas Mason Company, Inc., EPA-OHMSETT Facility, Leonardo, New Jersey) were obtained from the Directors Drive laboratory, labeled, and inserted into the routine on-site schedule for shipment to the laboratory for dioxin analysis.

4.1.3.6.3 Blank Spikes

Three dioxin-spiked wipes and three dioxin-spiked soils were issued concurrently with the blank samples described above.

Spiked wipe samples were prepared at the Directors Drive laboratory by soaking unused gauze wipes with solvent, then injecting 40 nanograms (ng) of dioxin directly onto each wipe. Samples were then individually labeled and inserted into the routine on-site schedule for shipment back to the laboratory for dioxin analysis.

Clarksburg soil spiked with 671 ppb of dioxin was obtained from the Directors Drive laboratory. The soil was spiked as part of a study performed for the USEPA-OHMSB by ITC (ITC, 1983). The dioxin level of 671 ppb was established by multiple analysis at the Directors Drive

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4.1.3.7 Corrective Action

Results for quality control check samples (in particular, field and trip blanks) were reviewed, as available, over the course of the project. Corrective actions were taken on an as-needed basis, including reanalysis of reextraction samples and resampling some locations when field blanks indicated potential contamination.

4.1.4 Analytical Methods

The methods used for the analysis of samples were either EPA or other approved analytical procedures. These methods were presented in detail in the Work Plan and significant modifications were not required during the analyses of the samples. Table 4.1.4-1 presents the matrices and the analytical parameters for which analysis was performed. Table 4.1.4-2 lists by parameter the analytes for each analysis.

As referred to in this report, priority pollutants are the organic base/neutral/acid (BNA) analysis, volatile organic compounds (VOC) analysis, pesticides and PCBs analysis, herbicide analysis, metals analysis, total cyanides, and total phenols. The BNA's, VOC's, and the pesticides and PCB's were analyzed according to the EPA Contract Laboratory Program (CLP) organic analysis requirements. Additionally, 40 extraneous peaks (as defined in the Work Plan) in the BNA and VOC analyses were library searched. All results were produced and reported according to the CLP protocols. The herbicides, metals, total cyanide, and total phenols were analyzed by EPA (1979, 1982) methods for waste analysis.

Dioxin was analyzed according to EPA protocols for dioxin in soils. Appropriate modifications were made for analyses of matrices other than soils. Detailed procedures are provided in the Work Plan. In addition to the analysis of dioxin, 10 percent of the soils (near surface and borings) and river sediments were analyzed for 2,3,7,8-tetra-chlorodibenzofuran and octachlorodibenzo-p-dioxin. Procedures for these methods are also provided in the Work Plan.

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- o Samples having concentrations sufficiently out of the linear calibration range would be reanalyzed as a one gram sample
- o One-gram samples that were still out of the linear calibration range would have their extracts diluted.

If reanalysis was required to obtain results within the linear calibration range, but insufficient or no sample remained after the initial analysis, only a dilution of the extract was performed.

4.2 SAMPLING, MONITORING, AND PHYSICAL TESTING

4.2.1 Ambient Air

Ambient air sampling was conducted at the site for establishing baseline conditions and for comparison of site data to data collected by the NJDEP at other sites in the Newark area.

A single air sampling location, the roof of the office/laboratory building, was utilized for the baseline sampling. This location (Figure 4.2.1-1) was approximately 10 meters above ground level and on the southern end of the site. Data from this single sampling location is considered representative of baseline ambient concentrations at the site because of the wind exposure at this point.

The samples and monitoring data were collected over 24-hour periods for 31 consecutive days beginning at noon on September 8, 1984, and ending at noon on October 9, 1984. Data were collected for a 31-day period so that an indication of the range of variation could be obtained.

The sampling and monitoring program included the measurement of the following parameters:

- o Total Suspended Particulate Matter
- o Inhalable Particulate Matter
- o Metals

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smaller than 10 microns turned with the air stream up into the impaction chamber and down through multiple vent tubes to the 8x10-inch glass fiber filter.

The high volume air sampler used was equipped with a constant flow control to maintain the 10-micron size selective inlet impactor cut-point, an elapsed-time indicator to accurately record sampling time period, and a pressure transducer flow recorder to continuously record pressure drop across the calibrated orifice. IPM concentration was computed by measuring the particulate mass collected and the total air volume sampled. Metals concentrations were computed by determining the total mass of individual metals collected and the total air volume sampled. Figure 4.2.1-3 is a schematic diagram the high-volume air sampler equipped with a 10-micron selective inlet.

A polyurethane foam (PUF) sampler was utilized to measure concentrations of dioxin, pesticides, and other chlorinated organics. Ambient air was drawn into a covered housing and through a dual-chambered aluminum sampling module. The upper chamber of the sampling module supported a four-inch diameter glass fiber filter used for collecting airborne particulates. The lower chamber consisted of an enclosed glass cartridge containing a 60-mm diameter by 75-mm-long cylindrical polyurethane foam plug for vapor entrapment. The sampling flow rate was approximately 0.25 cubic meter per minute. The PUF sampler was equipped with a calibrated venturi and magnehelic gage to measure flow rate, a voltage variator to adjust blower motor speed, and an elapsed-time indicator to accurately record sampling time period. Concentrations of dioxin, pesticides and other chlorinated organics were computed by measuring the total mass of each individual compound collected and the total air volume sampled. Figure 4.2.1-4 is a schematic diagram of the PUF sampler.

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Climatological data were also monitored continuously during the 31-day sampling period (noon, September 8, to noon, October 9, 1984). A Climatronics Mark III meteorological instrument for monitoring wind speed and direction was installed at the same location as the sampling instrumentation (Figure 4.2.1-1). Alignment of this instrument was checked daily. In addition to the site data, ambient temperature, barometric pressure, sky cover, precipitation, and wind speed and direction were obtained from National Weather Service Station WSO (Newark, New Jersey) which is located three miles southwest of the site.

4.2.2 Industrial Hygiene

The following industrial hygiene samples were taken during the field investigation:

- o Atmospheric Samples for Dioxin - A total of 67 samples were taken including 24 blanks. These samples were primarily personnel samples to define potential employee exposure and required respiratory protection. Some samples were taken in general work areas to determine background levels at various locations during work activities. To define potential respiratory protection breakthrough, the sampling tubes were spiked with ¹⁴C1-2,3,7,8-TCDD prior to sample collection.
- o Wipe Samples for Dioxin - A total of 14 wipe samples were taken, including three blanks. Most of these samples were taken to determine clean levels after equipment had been decontaminated. Four of the samples were taken in the employee decontamination area.
- o Water Sample for Dioxin - A water sample was taken from an employee decontamination pan.
- o Atmospheric Samples for Volatile Organics, Semi-volatile Organics, and Alcohols - A total of 12 samples were taken with charcoal tubes, fluorisil tubes, and silica gel tubes to identify atmospheric contaminant on site. Analysis for these samples was by gas chromatograph/mass spectrometry.

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The on-site buildings include four primary structures (Figure 4.0-1). The chemical manufacturing building with a 190-foot stack is located in the northeast corner of the property. The three-story process building is located west of the chemical manufacturing building along the edge of the Passaic River. A large warehouse/maintenance shop sits in the center of the site and a two-story, cement-block laboratory/office building is located near the entrance on the southern portion of the site.

In addition to the major buildings, a small solvent shed is located near the southwest corner of the property and a small pumphouse is situated between the process and warehouse buildings. There are also 142 tanks and vessels located throughout the site. Some are inside the various major buildings; others are outside, alone, or grouped together in the raw material or final product tank farms.

Three different types of samples were obtained to determine the level of dioxin contamination. Wipe samples were taken from painted or smooth surfaces where potentially contaminated dust or particulate had accumulated. Chip samples were collected from brick- and concrete-type surfaces that are porous or uneven and thus not suitable for wipe samples. Bulk samples were collected whenever significant amounts of dirt or dust existed in the buildings or when sufficient quantities of unknown liquids, solids, or sludges were present in the tanks or process vessels.

To obtain wipe samples, a 50-centimeter square template (2,500 centimeters squared area) was placed on a wall, floor, vessel side wall, etc. The area within the template was wiped with a three-by-three-inch sterile cotton gauze pad soaked with hexane. Sampling was conducted by applying pressure to the pad in straight, even strokes moving from left to right (horizontal for walls) in the area designated. The wipe effort was then repeated from top to bottom (vertical for walls) at a 90-degree

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areas of lower potential contamination such as the accounting office were not sampled.

The most probable sources of contamination on the vertical exterior walls of the building would be airborne particulates, dust from vehicle movement, and splatter from rainfall on contaminated ground. Because the building exterior consists of cement block and brick which are porous, chip samples were taken from the brick and concrete block surfaces. Exterior samples were taken from ground level to two feet above, from three to five feet above ground, and from the roof sill to two feet below at the center of the west and north walls and the southeast corner of the east wall. An additional sample was taken from the walkway of the main entrance to the building. These locations should be indicative of contamination caused by dust from vehicular traffic or by personnel contact with materials in the nearby drum storage areas. Because the roof of the office building is flat, contamination may have resulted from airborne particulates and from the air discharge from the laboratory hoods. A wipe sample was taken from the west end of the roof for analysis.

The types and numbers of samples collected from the interior and exterior of the office building are summarized in Table 4.2.3.2-1. Plan layouts of the first and second floors of the building with room identification numbers are shown in Figures 4.2.3.2-1 and 4.2.3.2-2, respectively.

Warehouse

The warehouse could have been contaminated from drum leakage and spillage, airborne particulates, and the tracking in of material on equipment and personnel. The warehouse is a steel frame building with corrugated cement panels forming the exterior walls and roof. The panels appear to be of asbestos-cement composition. The building has a pitched metal roof, and the entire structure is supported by a steel wall frame with

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Chip and wipe samples within the buildings were collected from concrete walls, floors, and columns near the process equipment. The sample locations were selected to represent the most likely areas of contamination such as the product packaging area and the acid conversion vessels. Bulk samples for dioxin analyses were collected from five locations in the process building. Sampling was conducted on all floors of both buildings.

Exterior samples from the building walls were taken in the same manner as the office/laboratory and warehouse buildings. The roof of the process building was wipe sampled in the northeast quadrant and the southwest corner. The roof of the chemical manufacturing building has collapsed into the structure, but two chip samples were collected from the top of the caved-in concrete slab.

A description of the samples collected from the process building and the chemical manufacturing building are provided in Tables 4.2.3.2-3 and 4.2.3.2-4, respectively.

Other Structures

The free-standing, 190-foot-high stack is adjacent to the southeast portion of the chemical manufacturing building. The stack is in poor structural condition, with small pieces of brick spalling from the outer surface. There are three openings in the stack. The lowest opening is a hinged manway at the base. A second opening is located where the flue enters the stack from the boiler, and the third is the top of the stack where smoke and gases exited. The two openings near the ground were sampled. A chip sample of soot was collected through the bottom manway. This sample should be a composite of material that has fallen from the interior of the stack since it was last in use. A chip sample was also taken from the interior of the flue pipe running from the boiler house to the stack. The exterior of the stack was sampled by taking a chip sample from ground level to two feet above.

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from building exterior corrugated panels. A total of 14 bulk samples were collected from the site--four from the office building, three from the warehouse, two from the chemical manufacturing building, three from the process building and two from process piping.

4.2.4 Sewers and Sumps

Four sewer and eight sump samples were collected from the site at the locations shown in Figure 4.2.4-1. Because existing plant drawings for the locations of these structures were limited, and the site was covered with geofabric, exterior sample locations were established with a metal detector. After a sump or sewer manhole cover was located, the geofabric was cut away. A visual inspection of the condition of the sump or sewer was made (i.e., depth of structure, presence of water, depth of sediment, etc.), and all information was recorded.

If the sump or sewer was dry, a sample was collected using a hand trowel and the material was placed in an aluminum pan. After sufficient material was collected for analytical testing, it was thoroughly mixed in the aluminum pan to produce a homogeneous sample and placed in sample bottles. At locations where the structure was too deep for collecting a sample by hand or where it contained water, a long-handled, perforated scoop was used for sample collection. Excess water was allowed to drain from the sample before it was placed in the aluminum pan for mixing and placing in sample bottles.

Data pertaining to sampling procedures, physical description of samples, amount collected, sample locations, etc., were recorded in the field log notebook. All sampling equipment, including trowels, gloves, pans, and scoops, was returned to the decontamination line for cleaning after the collection of each sample.

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strength before drilling operations continued. After the grout set, it was drilled out using an eight-inch roller bit and recycled drill water. Then sampling in the underlying silt layer was initiated.

The silt layer was sampled by continuous, undisturbed Shelby tube samples. Each Shelby tube was hydraulically pushed into the silt a maximum distance of two feet, allowed to stabilize in the borehole for 15 minutes, and then removed. After removal of the Shelby tube, the boring was advanced to the bottom depth of the previous Shelby tube sample with a four-inch roller bit and recirculating wash water. This drilling procedure was continued until the bottom of the silt layer was encountered. If sampling was to be performed below the silt layer, a four-inch PVC casing was installed to the depth of the silt/alluvial sand contact and tremie grouted to the ground surface. The casing was left undisturbed for at least 24 hours to allow proper curing of the grout mixture.

Sampling in the alluvial sands consisted of obtaining standard split-spoon samples at five-foot intervals and advancing the boring using rotary methods with water or drilling mud and a nominal four-inch roller bit (actual size was 3.75 inches). If field conditions warranted, a temporary steel casing was set in the boring to prevent caving of the borehole during drilling operations.

At the completion of a boring to its specified depth. The boring was tremie grouted to the surface using cement grout as the temporary steel casing was removed. All water forced to the surface during the grouting procedure was collected and stored on site.

4.2.3.1.2 Sample Collection Procedures

At each boring location, the first sample was taken from depths of zero to six inches using a hand trowel. The sample was placed in an aluminum

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DEPTH	ANALYSIS AND SAMPLE CONTAINER
0 - 6 inches	Full priority pollutants and dioxin; three 500-ml jars
6 - 12 inches	Dioxin; one 250-ml jar
12 - 24 inches	Full priority pollutants and dioxin; three 500-ml jars
24 inches below the surface to 18 inches above the bottom of fill	Selected dioxin; one 250-ml jar
18 inches above the bottom of fill to 6 inches above bottom of fill	Full priority pollutants; three 500-ml jars
6 inches above bottom of fill to the bottom of the fill	Dioxin; one 250-ml jar

Samples of 250 milliliters were taken from the top and bottom of each Shelby tube and composited. The second Shelby tube sample from the silt layer in each boring was analyzed for dioxin. When a dioxin level greater than one part per billion (ppb) was observed, selected Shelby tube samples from the same boring were also tested for dioxin (remaining tubes were archived).

4.2.5.2 Near-Surface Soil Sampling

Near-surface soil sampling locations were chosen using a biased approach based on plant activities and functional units. Sample location points are shown in Figure 4.2.5-1. Samples were obtained from depths of zero to 6 inches, 6 to 12 inches, and 12 to 24 inches at each location, with additional samples collected from 24 to 36, 36 to 48, and 48 to 60 inches wherever possible. Each near-surface soil sampling location was identified by its grid coordinate system location, as previously described.

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Samples collected from zero to 6 inches and 6 to 12 inches were obtained using a hand trowel or by hand. Material collected from the entire sampling interval was placed in an aluminum mixing pan and thoroughly composited before being placed into sample bottles. Samples were placed in either 250- or 500-milliliter amber sampling bottles depending on the analysis to be performed.

After completion of sampling from the upper 12 inches in unconsolidated materials, the hole was cased with an eight-inch PVC pipe. The eight-inch casing was grouted in the hole with a quick-drying hydraulic cement to anchor it and prevent cross contamination of the lower sampling increment from materials sloughing down the outside.

Samples collected from 12 to 24 inches were obtained with a post-hole digger, hand auger, hand trowel, or by hand. At some locations where brick or concrete debris was encountered, a steel digging bar was used to break up or loosen the material to be sampled. Material collected from the entire sample length was placed in an aluminum mixing pan and composited. Upon completion of sampling to 24 inches, a four-inch PVC casing was installed with either cement grout or quick-drying hydraulic cement before additional samples were collected. At locations where cement grout was used, the grout was allowed to cure for at least 24 hours before additional samples were collected.

If a sampling location was in an area covered by concrete or asphalt, an eight-inch PVC casing was installed from the top of the slab to a depth 12 inches below. Grout was not necessary to hold this casing in place. After obtaining the sample from 12 to 24 inches, a four-inch PVC casing was installed using a neat concrete grout. The grout was allowed to cure for a minimum of 24 hours before additional samples were obtained.

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4.2.6.1 Well Installation

Shallow Monitoring Wells

Shallow monitoring wells were located within 10 feet of the corresponding soil boring. The well boreholes were advanced with 12-inch hollow-stem augers to the top of the silt layer. A schematic diagram of a typical shallow monitoring well is shown in Figure 4.2.6.1-1.

For the shallow monitoring wells, Schedule 40, flush coupled and jointed, two-inch-diameter PVC casing with unglued slip caps on both the top and the bottom of the casing was used. The screen was generally 5 to less than 10 feet (as per NJDEP screening requirements) of No. 10 slot (0.010-inch) two-inch-diameter Schedule 40 PVC. The top of the screen was located approximately one foot above the static water table. The filter pack consisted of clean bagged sand approved by NJDEP for use on site.

A layer of bentonite pellets (0.2 to 0.5 foot thick) was placed at the top of the gravel pack to prevent penetration of the gravel pack by the grout. This layer was tamped after placement to provide an effective seal. The remainder of the annular space was filled with cement grout to minimize surface infiltration. All monitoring wells were grouted prior to the end of a shift, and a six-inch diameter protective steel outer casing was installed following the completion of each well. A protective concrete collar 1-1/2 feet square and a minimum of one foot thick was constructed at or near the ground surface around the protective casing.

Intermediate Monitoring Well

An intermediate depth monitoring well was located approximately 100 feet south of the site, as shown in Figure 4.2.6-1. The well was screened in the alluvial sand which underlies the silt at the site. To reduce the potential for surface contaminants being introduced into the alluvial

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screen as the casing was withdrawn. A tremie pipe was set in the annular space outside of the casing to the bottom of the silt and grout was introduced, flushing the mud out as the well was grouted to the surface.

4.2.6.2 Well Development

Following installation of the well, development was performed to remove materials introduced into the screened soil strata during drilling. All wells were developed for a minimum period of one hour or until a turbidity-free discharge was obtained. Surging was also employed for wells installed in the upper fine-grained fill materials. Well development was performed at least two days after well completion to minimize the potential for drawing uncured grout into the filter pack of the well. All water resulting from well development was collected and stored on site.

4.2.6.3 Water Level Monitoring

Ground water level measurements were taken in the eight perimeter monitoring wells at approximately 40-minute intervals for a 12-hour period on October 15, 1984. At the same time, tidal measurements were taken at the staff gage located on the river close to monitoring well MW-2A. The staff gage measurements were taken to ensure that a complete tidal cycle was covered. Continuous-reading water level recorders were installed in monitoring wells MW-1A and MW-7A to evaluate the hydraulic connection between the Passaic River and the water-bearing zones beneath the site.

4.2.6.4 Ground Water Sampling

Following the installation and development of the monitoring wells, ground water sampling was conducted on October 9 and October 30, 1984, to assess the extent of ground water contamination. Each set of ground water samples consisted of one sample from each well. The procedure for obtaining ground water samples is described in the following paragraphs.

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4.2.6.5 Slug Tests

Slug tests were conducted in monitoring wells MW-1A through MW-8A on November 19 and 20, 1984, to estimate the hydraulic conductivity (permeability) of the zones screened by each well. The test consisted of measuring the rate at which the water level in the monitoring well declined after a known volume (slug) was rapidly introduced and the rate at which the water level rose after the slug was rapidly removed. Two slugs were used at the site--a three-foot-long PVC slug in monitoring wells MW-3A through MW-8A and a four-foot-long solid steel slug in monitoring wells MW-1A and MW-2A. The volumes of the slugs were small compared to the volume of water in the aquifer; thus, the test provided an estimate of hydraulic conductivity within only a few feet radius of each monitoring well. An ENVIROLABS Model DL-120-MCP Data-Logger with a submersible pressure transducer was used to record water level changes.

The slug was quickly, but smoothly, lowered into the well until immersed. The water level initially rose as the slug was immersed and then fell to reach equilibrium with the aquifer water level. With the exception of monitoring wells MW-2A and MW-6A, the resulting falling water level was recorded on the strip chart until the water level returned to its pre-immersion static level. Then the slug was quickly and smoothly removed from the well. The resulting rise in water level was recorded on the strip chart. When the water level returned to static level, the test was considered complete.

4.2.7 Passaic River Water

4.2.7.1 River Level Monitoring

A staff gage (Figure 4.2.7.1-1) was installed in the Passaic River adjacent to the site near monitoring well MW-2A, and the tidal fluctuations in the river level were measured through a full tidal cycle on two occasions. The observed levels were referenced to the site datum. Results of these measurements on October 15, 1984, were compared to

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site, 300 feet upstream from this transect, and 100 feet downstream from the east property line. Figure 4.2.1-8 indicates the locations of the samples.

Samples were taken from three locations along each of the three transects. These locations were the north bank, center, and south bank of the Passaic River. The samples collected from the transect locations were from depth intervals of zero to 12 inches and 12 to 24 inches. Samples from zero to 12 inches were obtained from the remaining 14 sample locations, and four of the remaining locations on the south bank were sampled from 12 to 24 inches.

River sediment samples were collected by advancing Shelby tubes to the prescribed sampling depth and recovering a sediment core sample. The sample was extruded into a clean aluminum pan and the edges scraped with a stainless steel spoon to minimize cross contamination. Only the central core of the sediment sample was transferred to a sample bottle for chemical analysis. If samples were required from two depths (zero to 12 inches and 12 to 24 inches), the sediment column from each interval was extruded into a clean aluminum pan. All sampling equipment was decontaminated between sample collections.

Following the completion of the surface sediment sampling, a deep core sediment sample was taken at Station 1-3-0. A PVC casing was pushed into the sediment to a depth of 40 inches. The sediments were compressed by the casing, however, and only six inches of material were recovered. The remaining core samples were obtained using a piston sampler, which was decontaminated between samples. Sample increment depths were: 40 to 46 inches, 46 to 52 inches, 60 to 66 inches, and 66 to 72 inches.

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Sampling of the drums was conducted by a four-man crew consisting of a foreman and three technicians. The foreman supervised and recorded data in the field log notebook. Self-contained breathing apparatus was worn by Technicians 1 and 2 during the drum opening and sampling activities; the foreman and Technician 3 wore full-faced respirators. Technician 1 wore a heavy PVC suit; the other members of the sampling team wore polyethylene-coated tyvek suits. Technician 1 opened the drums and sealed them after sampling, and performed air monitoring for organic vapors and combustible gases during drum sampling. Technician 2 sampled the drums and performed air monitoring during opening and sealing of the drum. Technician 3 changed air cylinders for the breathing apparatus, returned the samples to the laboratory trailer, and stenciled the drums.

All drums were sampled on a "where-is" basis with only minimal movement to permit access to the drum. Prior to and during drum opening activities, monitoring was conducted using a HNU/PID organic vapor detector and a Castex 3 (CX-3) combustible gas indicator.

The bungs or open heads were removed using nonsparking tools. Only one drum at a time was opened to minimize the release of any combustible or organic vapor. Glass sampling rods (approximately 3 feet by 13 millimeters) were used to sample liquids. The rods were inserted to the bottom of the drum, the exposed end of the glass rod was plugged, and the sample was transferred to the sample bottle. The rod was then disposed of in the drum and the drum was resealed. Electrical conduit was used in a manner similar to that described above to sample drums containing friable or crystallized material. Drums containing extremely hard or solid material were sampled using a stainless steel boring tool. The tool was decontaminated after each use.

Sampling data recorded in the field log notebook by the foreman included date, time, type of drum, description of the sample, percent LEL reading, and the organic vapor level reading in parts per million.

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TABLES

TABLE 4.1.3.2-1
SAMPLE PACKAGING REQUIREMENTS

HAZARDOUS (POISON B PACKAGING)	NONHAZARDOUS (CONVENTIONAL PACKAGING, PRESERVATION IN ICE CHESTS)
Site Soils (near-surface and borings)	Site Waters
Wipes	Industrial Hygiene
Drums	Ambient Air
Tanks	River Sediments

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TABLE 4.1.3.2-2
SAMPLE PRESERVATION REQUIREMENTS

PARAMETER	TECHNIQUE	
	SOIL	WATER
Dioxin	None	None
Organic PP	Cool, 4°C ⁽¹⁾	Cool, 4°C ⁽¹⁾
Metals	None	2 ml conc. HNO ₃ ⁽²⁾ (to pH <2)
Cyanide	None	2 ml conc. NaOH ⁽²⁾ (to pH >12)
Phenols	None	2 ml conc. H ₂ SO ₄ ⁽²⁾ (to pH <2)

(1) When possible, with packaging restrictions.

(2) Added to sample bottles prior to collection; these containers must not be rinsed prior to being filled with sample.

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TABLE 4.1.3.5-1
ROUTINE QUALITY CONTROL SAMPLES AND
LABORATORY CHECK FREQUENCIES

	FIELD BLANKS	TRIP BLANKS	ON-SITE SPLITS WITH NJDEP	METHOD BLANKS	BLIND SPLITS	SAMPLE OR BLANK SPIKES	INTERNAL SURROGATE STANDARDS	REFERENCE (CALIBRATION STANDARDS)
DIOXIN								
Wipes	5	5	-	5	-	5	100	1/shift
Soil/Sediment/Sludge	-	-	5	5	5	5	100	1/shift
Water	5	5	5	5	5	5	100	1/shift
Chips	-	-	5	5	5	5	100	1/shift
VOLATILES								
Soil/Sediment/Sludge	5	5	5	5	5	5	100	1/shift
Water	5	5	5	5	5	5	100	1/shift
SEMI-VOLATILES								
Soil/Sediment/Sludge	-	-	5	5	5	5	100	1/shift
Water	5	5	5	5	5	5	100	1/shift
PP METALS								
Soil/Sediment/Sludge	-	-	5	5	5	5	-	1/shift
Water	5	5	5	5	5	5	-	1/shift
CYANIDE								
Soil/Sediment/Sludge	-	-	5	5	5	5	-	1/shift
Water	5	5	5	5	5	5	-	1/shift
TOTAL PHENOLS								
Soil/Sediment/Sludge	-	-	5	5	5	5	-	1/shift
Water	5	5	5	5	5	5	-	1/shift

Note: All numbers are percentages.

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TABLE 4.1.4-1
ANALYSIS PARAMETERS VERSUS SAMPLE MATRICES

ANALYTICAL PARAMETERS	NUMBER OF PARAMETERS OR ANALYTES/ ANALYTICAL METHOD	SAMPLE TYPE	SOIL	RIVER SEDI- MENTS	SEWER SLUDGES	WELL BORINGS (SOIL)	AIR	WELL WATER	DROPS	WIPES	CHIPS	SCRAPES	RIPIRATE WATER	BULK	INDUS- TRIAL HYGIENE
2,3,7,8-TCDD	(1)		X	X	X	X	X	X	X	X	X	X	X		X
2,3,7,8-TCDF ^a	(1)		X	X		X									
Octachlorodioxin ^a	(1)		X	X		X									
Priority Pollutant Acid Base/Neutrals (Ac/B/N)	(69)		X	X		X		X					X		
Priority Pollutant Pesti- cides	(25)		X	X		X		X					X		
Priority Pollutant Metals	(13)		X	X		X		X					X		
Priority Pollutants Volatile Organic Compounds (VOC)	(30)		X	X		X		X					X		
Herbicides	(10)		X	X		X		X					X		
Polycyclic Aromatic Hydrocarbons (PAH)	(25)						X								
Ambient Air Volatile Organic Compounds	(24)						X								
Asbestos	(1)						X					X		X	X
Ambient Air Metals	(8)						X								
Inhalable Particulate	(1)						X								
Total Suspended Solids	(1)						X								
Vinyl Chloride	(1)						X								
Pesticides and Other Chlorinated Organics	(7)						X								
Hazardous Waste Charac- terization	-								X						
Cyanides	(1)		X	X		X		X					X		
Phenols	(1)		X	X		X		X					X		
Others															X

^a 10 percent of total samples collected.

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TABLE 4.1.4-2
LISTING OF ANALYTES

CAS NUMBER	COMPOUND NAME
<u>DIOXIN</u>	
1746-01-6	2,3,7,8-Tetrachloro- dibenzo-p-dioxin
	2,3,7,8-Tetrachloro- dibenzofuran
3268-87-9	Octachlorodibenzo-p- dioxin
<u>PRIORITY POLLUTANT</u>	
<u>Volatile Organic Compounds</u>	
71-43-2	Benzene
56-23-5	Carbon tetrachloride
108-90-7	Chlorobenzene
107-06-2	1,2-Dichloroethane
71-55-6	1,1,1-Trichloroethane
75-34-3	1,1-Dichloroethane
79-00-5	1,1,2-Trichloroethane
79-34-5	1,1,2,2-Tetrachloroethane
75-00-3	Chloroethane
542-88-1	Bis(chloromethyl) ether
110-75-8	2-Chloroethylvinyl ether
67-66-3	Chloroform
75-35-4	1,1-Dichloroethene
156-60-5	trans-1,2-Dichloroethene
78-87-5	1,2-Dichloropropane
10061-02-6	trans-1,3-Dichloro-propene
10061-01-5	cis-1,3-Dichloro-propene
100-41-4	Ethylbenzene
75-09-2	Methylene chloride
74-87-3	Chloromethane
74-83-9	Bromomethane
75-25-2	Bromoform
75-27-4	Bromodichloromethane
75-69-4	Trichlorofluoromethane
75-71-8	Dichlorodifluoromethane
124-48-1	Chlorodibromomethane
127-18-4	Tetrachloroethene
108-88-3	Toluene
79-01-6	Trichloroethene
75-01-4	Vinyl chloride

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TABLE 4.1.4-2
(Continued)

CAS NUMBER	COMPOUND NAME
67-64-1	Acetone
78-93-3	2-Butanone
75-15-0	Carbon disulfide
519-78-6	2-Hexanone
108-10-1	4-Methyl-2-pentanone
100-42-5	Styrene
108-05-4	Vinyl acetate
95-47-6	Total Xylenes

PRIORITY POLLUTANT

Base/Neutral and Acid Organic Compounds

88-06-2	2,4,6-Trichlorophenol
59-50-7	4-Chloro-3-methyl-phenol
95-57-8	2-Chlorophenol
120-33-2	2,4-Dichlorophenol
105-67-9	2,4-Dimethylphenol
88-75-5	2-Nitrophenol
100-02-7	4-Nitrophenol
51-28-5	2,4-Dinitrophenol
534-52-1	4,6-Dinitro-2-methylphenol
87-86-5	Pentachlorophenol
108-95-2	Phenol
65-85-0	Benzoic acid
95-48-7	2-Methylphenol
108-39-4	4-Methylphenol
95-95-4	2,4,5-Trichlorophenol
83-32-9	Acenaphthene
92-87-5	Benzidine
120-82-1	1,2,4-Trichlorobenzene
118-74-1	Hexachlorobenzene
67-72-1	Hexachlorethane
111-44-4	Bis(2-chloroethyl)ether
91-58-7	2-Chloronaphthalene
95-50-1	1,2-Dichlorobenzene
541-73-1	1,3-Dichlorobenzene
106-46-7	1,4-Dichlorobenzene
91-94-1	3,3'-Dichlorobenzidine
121-14-2	2,4-Dinitrotoluene
606-20-2	2,6-Dinitrotoluene
122-66-7	1,2-Diphenylhydrazine
206-44-0	Fluoranthene
7005-72-3	4-Chlorophenyl phenyl ether
101-55-3	4-Bromophenyl phenyl ether
39638-32-9	Bis(2-chloroisopropyl)ether
111-91-1	Bis(2-chloroethoxy)methane
87-68-3	Hexachlorobutadiene
77-47-4	Hexachlorocyclopentadiene
78-59-1	Isophorone

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TABLE 4.1.4-2
(Continued)

CAS NUMBER	COMPOUND NAME
91-20-3	Naphthalene
98-95-3	Nitrobenzene
62-75-9	N-nitrosodimethylamine
86-30-6	N-nitrosodiphenylamine
621-64-7	N-nitrosodipropylamine
117-81-7	Bis(2-3ethylhexyl)phthalate
85-68-7	Butyl benzyl phthalate
84-74-2	Di-N-butyl phthalate
117-84-0	Di-N-octyl phthalate
84-66-2	Diethyl phthalate
131-11-3	Dimethyl phthalate
56-55-3	Benzo(A)anthracene
50-32-8	Benzo(A)pyrene
205-99-2	Benzo(B)fluoranthene
207-08-9	Benzo(K)fluoranthene
218-01-9	Chrysene
208-96-8	Acenaphthylene
120-12-7	Anthracene
191-24-2	Benzo(GHI)perylene
86-73-7	Fluorene
85-01-0	Phenanthrene
53-70-3	Dibenzo(A,H) anthracene
193-39-5	Indeno(1,2,3-CD)pyrene
129-00-0	Pyrene
62-53-3	Aniline
100-51-6	Benzyl alcohol
106-47-8	4-Chloroaniline
132-64-9	Dibenzofuran
91-57-6	2-Methylnaphthalene
88-74-4	2-Nitroaniline
99-09-2	3-Nitroaniline
100-01-6	4-Nitroaniline

PRIORITY POLLUTANT

Pesticides and PCBs

309-00-2	Aldrin
60-57-1	Dieldrin
57-74-9	Chlordane
50-29-3	4,4'-DDT
72-55-9	4,4'-DDE
72-54-8	4,4'-DDD
959-98-8	alpha-Endosulfan
33213-65-9	beta-Endosulfan
1031-07-8	Endosulfan sulfate
72-20-8	Endrin
7421-93-4	Endrin aldehyde
76-44-8	Heptachlor
1024-57-3	Heptachlor epoxide

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TABLE 4.1.4-2
(Continued)

CAS NUMBER	COMPOUND NAME
319-84-6	alpha-BHC
319-85-7	beta-BHC
58-89-9	gamma-BHC
319-86-8	delta-BHC
53469-21-9	PCB-1242
11097-69-1	PCB-1254
11104-28-2	PCB-1221
11141-16-5	PCB-1232
12672-29-6	PCB-1248
11096-82-5	PCB-1260
12674-11-2	PCB-1016
8001-35-2	Toxaphene

PRIORITY POLLUTANT

Chlorinated Herbicides

75-99-0	Dalapon (Dowpon)
1918-00-9	Dicamba
7085-19-0	MCPP
94-74-6	MCPA
120-36-5	Dichloroprop (2,4-DP)
94-75-7	2,4-D
93-72-1	2,4,5-TP (Silvex)
93-76-5	2,4,5-T
94-82-6	2,4-DB
88-85-7	Dinoseb (DNBP)

PRIORITY POLLUTANT

Metals

Antimony
Arsenic
Beryllium
Cadmium
Chromium
Copper
Lead
Mercury
Nickel
Selenium
Silver
Thallium
Zinc

TABLE 4.1.4-2
(Continued)

CAS NUMBER	COMPOUND NAME
------------	---------------

Classical Parameters

	Total Cyanide
	Total Phenols
	Asbestos

AMBIENT AIR

Metals

	Lead
	Manganese
	Copper
	Vanadium
	Cadmium
	Zinc
	Iron
	Nickel

AMBIENT AIR

Volatile Organic Compounds (VOCs)

	Vinyl chloride
	Vinylidene chloride
	Methylene chloride (ME chloride)
	Chloroform
	1,2-Dichloroethane
	Benzene
	Carbon tetrachloride
	Trichloroethylene (TRIC)
	1,4-Dioxane
	1,1,2-Trichloroethane
	toluene
	1,2-Dibromomethane
	Tetrachloroethylene (PERC)
	Chlorobenzene
	Ethylbenzene
	m-Xylene
	p-Xylene
	Styrene
	o-Xylene
	1,1,2,2-Tetrachloroethane
	o-Chlorotoluene
	p-Chlorotoluene
	p-Dichlorobenzene
	o-Dichlorobenzene
	Nitrobenzene

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TABLE 4.1.4-2
(Continued)

CAS NUMBER	COMPOUND NAME
------------	---------------

AMBIENT AIR

Polycyclic Aromatic Hydrocarbons (PAHs)

	Benzo(k)fluoranthene
	Benzo(a)pyrene
	Benzo(ghi)perylene
	Indeno(1,2,3-cd)pyrene
	Coronene
	Phenanthrene
	Triphenylene
	Benzo(b)fluoranthene
	Anthracene
	Fluoranthene
	Pyrene
	Benz(a)anthracene
	Benz(ah)anthracene
	Chrysene
	Perylene

AMBIENT AIR

Pesticides and Other Chlorinated Organics

	Benzene sulfonyl chloride
	Tetrachlorobenzene
	4-chlorobenzene sulfonyl chloride
	4-methoxybenzene sulfonyl chloride
	Hexachlorobenzene
	2,4,5-T (methyl ester)
	Ovex
	p,p'-DDT
	Total suspended particulate
	Total inhalable particulate

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TABLE 4.2.1-1
AMBIENT AIR ANALYSIS COMPONENTS

I. METALS

Lead	Cadmium
Manganese	Zinc
Copper	Iron
Vanadium	Nickel

II. VOLATILE ORGANIC COMPOUNDS (VOC's)

Vinyl chloride	Tetrachloroethylene (PERC)
Vinylidene chloride	Chlorobenzene
Methylene chloride (ME chloride)	Ethylbenzene
Chloroform	m-Xylene
1,2-Dichloroethane	p-Xylene
Benzene	Styrene
Carbon tetrachloride	o-Xylene
Trichloroethylene (TRIC)	1,1,2,2-Tetrachloroethane
1,4-Dioxane	o-Chlorotoluene
1,1,2-Trichloroethane	p-Chlorotoluene
Toluene	p-Dichlorobenzene
1,2-Dibromoethane	o-Dichlorobenzene
Nitrobenzene	

III. POLYCYCLIC AROMATIC HYDROCARBONS (PAH's)

Benzo(k)fluoranthene	Anthracene
Benzo(a)pyrene	Fluoranthene
Benzo(ghi)perylene	Pyrene
Indeno(1,2,3-cd)pyrene	Benz(a)anthracene
Coronene	Benz(ah)anthracene
Phenanthrene	Chrysene
Triphenylene	Perylene
Benzo(b)fluoranthene	

IV. PESTICIDES AND OTHER CHLORINATED ORGANICS

Benzene sulfonyl chloride
Tetrachlorobenzene
4-Chlorobenzene sulfonyl chloride
4-Methoxybenzene sulfonyl chloride
Hexachlorobenzene
2,4,5-T (methyl ester)
Oxex
p,p'-DDT

V. ASBESTOS

VI. TETRACHLORODIBENZO DIOXIN (2,3,7,8-TCDD)

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TABLE 4.2.3-1
SAMPLE SUMMARY FOR BUILDINGS,
STRUCTURES, AND EQUIPMENT

DESCRIPTION		WIPE	CHIP	BULK
Office Building				
Interior	Areas of first and highest exposure (lab, change room, etc.)	22	5	4 (asbestos)
Exterior	Primary contact areas near roads and sidewalks	1	10	-
Warehouse				
Interior	Areas of highest exposure (office, lunchroom, shop)	7	4	3 (asbestos)
Exterior	Contact areas (splashing and wind borne)	1	12	-
Manufacturing Building				
Interior	Packaging and reaction areas	4	14	2 (asbestos)
Exterior	Wind-borne contact	1	9	-
Process Building				
Interior	Reaction and carbon infiltration areas	12	3	3 (asbestos) 5 (dioxin)
Exterior	High source area (example, carbon filter)	2	7	-
Stack	Flue, sludge pit, outer surface	-	3	-
Solvent Storage Shed	Representative internal and external	-	1	1 (dioxin)
Well House	Representative internal and external	-	2	-
Tanks (in buildings tank and farms)		28	-	112 (dioxin) 2 (asbestos)
TOTAL		78	70	132

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TABLE 4.2.3.2-1

OFFICE AND LABORATORY BUILDING SAMPLE LOCATIONS

SAMPLE NUMBER	DESCRIPTION
<u>WIPE SAMPLES</u>	
1100-0016-W-L	Lab Room 1100, Main Entrance
1102-0017-W-L	Lab Room 1102, Accounting
1105-0018-W-L	Lab Room 1105, Floor, Plant Manager
1106-0021-W-L	Lab Room 1106, Floor, Back Foyer Inside Door
1107-0020-W-L	Lab Room 1107, Floor
1108-0019-W-L	Lab Room 1108, Wall
1116-0034-W-L	Lab Room 1116, Locker Room
1122-0035-W-L	Lab Room 1122, Heater Duct, Basket Room
1122-0073-W-L	Lab Room 1122, Windowsill, Basket Room
1122-0074-W-L	Lab Room 1122, Floor Near Inside Entrance
1202-0032-W-L	Lab Room 1202, Floor, Lunchroom
1202-0033-W-L	Lab Room 1202, Radiator, Lunchroom
1204-0023-W-L	Lab Room 1204, Floor by Back Door, Lab
1204-0024-W-L	Lab Room 1204, Lab Hood, Lab
1204-0025-W-L	Lab Room 1204, North Side of Entrance, Lab Side
1204-0026-W-L	Lab Room 1204, Bench Near Back Door
1205-0030-W-L	Lab room 1205, A/C Intake Duct, Utility Room
1205-0031-W-L	Lab Room 1205, Furnace Intake, Utility Room
1205-0095-W-L	Lab Room 1205, Heater Interior Inlet, Utility
1206-0027-W-L	Lab Room 1206, Floor, Small Lab
1206-0028-W-L	Lab Room 1206, Bench, Small Lab
1206-0381-W-L	Lab Room 1206, Bench, Small Lab
1506-1590-W-L	Office/Lab, West Wall, at Roof

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TABLE 4.2.3.2-1
(Continued)

SAMPLE NUMBER	DESCRIPTION
<u>CHIP SAMPLES</u>	
1118-0049-C-L	Lab Room 1118, Floor Under Sink Edge, Washroom
1119-0050-C-L	Lab Room 1119, Floor Slop Sink
1122-0051-C-L	Lab Room 1122, Floor Under Arch Between Room 1122 and 1116
1122-0052-C-L	Lab Room 1122, Floor Near Drain, Basket Room
1122-0053-C-L	Lab Room 1122, Floor Near Back Door, Basket Room
1501-0098-C-L	Lab Exterior, 1501, Center, North Wall at Roof Sill
1501-0111-C-L	Lab Exterior, 1501, Center of North Wall, 3 to 5 feet
1501-0113-C-L	Lab Exterior, 1501, Center North Wall, Ground Level
1505-0097-C-L	Lab Exterior, 1505, South Corner, East Wall at Roofsill
1505-0108-C-L	Lab Exterior, 1505, South Corner, East Wall, 3 to 5 feet
1505-0109-C-L	Lab Exterior, 1505, South Corner, East Wall, Ground Level
1505-0110-C-L	Lab Exterior, 1505, Walkway of Front Entrance
1506-0099-C-L	Lab Exterior, 1506, Center West Wall, Top 24-inch Vertical
1506-0166-C-L	Lab Exterior, 1506, Center West Wall, 3 to 5 feet
1506-0167-C-L	Lab Exterior, 1506, Center West Wall, Ground Level

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TABLE 4.2.3.2-2
WAREHOUSE SAMPLE LOCATIONS

SAMPLE NUMBER	DESCRIPTION
<u>WIPE SAMPLES</u>	
2100-0218-W-L	Warehouse, Room 2100, Top of Fluorescent Light
2103-0217-W-L	Warehouse, Room 2103, Floor, Foreman's Office
2108-0176-W-L	Warehouse, Room 2108, Floor, Kitchen
2108-0177-W-L	Warehouse, Room 2108, Window Sill, Kitchen
2109-0178-W-L	Warehouse, Room 2109, Top of Light Work Area, Shop
2109-0179-W-L	Warehouse, Room 2109, Top of Bench in Shop
2200-0180-W-L	Warehouse, Room 2200, Top of Beam in Storage Area
2400-0315-W-L	Warehouse, West End, Roof
<u>CHIP SAMPLES</u>	
2100-0168-C-L	Warehouse, Room 2100, Center of Traffic Area, Floor
2109-0169-C-L	Warehouse, Room 2109, Floor, Tool Crib Cage Area
2109-0170-C-L	Warehouse, Room 2109, Floor by Traffic Door
2109-0171-C-L	Warehouse, Room 2109, Floor by Warehouse Door
2501-0317-C-L	Warehouse, North Wall at Ground Level
2501-0391-C-L	Warehouse, North Wall at 60 inches (3 to 5 feet)

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Warehouse, Exterior, North Side at Roof	2501-0392-C-L
Warehouse, South Wall at Ground Level	2502-0319-C-L
Warehouse, South Wall at 60 inches (3 to 5 feet)	2502-0393-C-L
Warehouse, Exterior, South Wall, at Roof Line	2502-0529-C-L
Warehouse, East Wall at Ground Level	2504-0318-C-L
Warehouse, Exterior, East Wall, 3 to 5 feet	2504-0527-C-L
Warehouse, Exterior, East Wall, at Roof Line	2504-0528-C-L
Warehouse, West Wall at Ground Level	2506-0316-C-L
Warehouse, West Wall at 60 inches (3 to 5 feet)	2506-0389-C-L
Warehouse, West Wall at Roof Line	2506-0390-C-L

CHIP SAMPLES

SAMPLE NUMBER	DESCRIPTION
---------------	-------------

TABLE 4.2.3.2-2
(Continued)

TABLE 4.2.3.2-3
PROCESS BUILDING SAMPLE LOCATIONS

SAMPLE NUMBER	DESCRIPTION
<u>WIPE SAMPLES</u>	
4100-0556-W-L	Process Bldg., East End, Near Vessel, Top of Light, First Floor
4100-0557-W-L	Process Bldg., East End, Low on Column, Near Vessel, First Floor
4100-0558-W-L	Process Bldg., Center, First Floor, Top of Light, Near Vessel
4100-0559-W-L	Process Bldg., Center, First Floor, Low on Column, Near Vessel
4100-0560-W-L	Process Bldg., West End, First Floor, Top of Light, Near Vessel
4100-0561-W-L	Process Bldg., West End, First Floor, Low on Column, Near Vessel
4200-0608-W-L	Process Bldg., Second Floor, West End Interior Wall
4200-0609-W-L	Process Bldg., Second Floor, Acid Room Wall (Interior)
4200-0610-W-L	Process Bldg., Second Floor, East End Interior Wall
4300-0611-W-L	Process Bldg., Third Floor, East End Interior Wall
4300-0612-W-L	Process Bldg., Third Floor, Surface, Center
4300-0613-W-L	Process Bldg., Third Floor, Surface, East End
4400-0495-W-L	Process Bldg., Roof, Northeast Quadrant
4400-0496-W-L	Process Bldg., Roof, Southwest Corner
<u>CHIP SAMPLES</u>	
4100-0553-C-L	Process Bldg., Floor, West End of First Floor
4100-0554-C-L	Process Bldg., Floor at Loading Door, First Floor
4100-0555-C-L	Process Bldg., Floor, East End Under Vessel, First Floor
4501-0424-C-L	Process Bldg., Exterior, North Wall, 0 to 24 inches
4502-0451-C-L	Process Bldg., South Wall, Near Roof at Vert Stairs, Exterior
4503-0427-C-L	Process Bldg., Exterior, South at C Filter, 24 inches Over Curb

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TABLE 4.2.3.2-3
(Continued)

SAMPLE NUMBER	DESCRIPTION
<u>CHIP SAMPLES</u>	
4504-0452-C-L	Process Bldg., East Wall, Over Trench Near Vessels (0 to 24 inches)
4506-0425-C-L	• Process Bldg., Exterior Bin Wall, West Side, 0 to 24 inches
4506-0426-C-L	Process Bldg., Exterior, Bin Wall, West Side, 36 to 60 inches
<u>BULK SAMPLES</u>	
4501-0455-B-L	Process Bldg., North Wall, 36 to 60 inches
4501-0493-B-L	Process Bldg., North Wall, 24 inches from top (Off Louvers)
4503-0456-B-L	Process Bldg., South Wall, 36 to 60 inches
4504-0453-B-L	Process Bldg., East Wall, Near Vessels (36 to 60 inches)
4504-0454-B-L	Process Bldg., East Wall, at Roof Near Vessels

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TABLE 4.2.3.2-4
CHEMICAL MANUFACTURING BUILDING SAMPLE LOCATIONS

SAMPLE NUMBER	DESCRIPTION
<u>WIPE SAMPLES</u>	
3100-0658-W-L	Mftg. Bldg., First Floor, Packing Area, Rafter
3200-0655-W-L	Mftg. Bldg., Second Floor, New Addition, Floor, South End
3200-0656-W-L	Mftg. Bldg., Second Floor, New Addition, Panel, Center
3200-0657-W-L	Mftg. Bldg., Second Floor, New Addition, North End, Beam
3502-0716-W-L	Mftg. Bldg., South Exterior Door
<u>CHIP SAMPLES</u>	
3100-0619-C-L	Mftg. Bldg., Old Area, Roof Slab, South of Center Vessel
3100-0620-C-L	Mftg. Bldg., Old Area, Roof Slab, West of North Vessel
3100-0621-C-L	Mftg. Bldg., Bulk Debris from Drain Area
3100-0622-C-L	Mftg. Bldg., Old Area, 1st Floor, Floor North End North Room
3100-0633-C-L	Mftg. Bldg., Old Area, Floor, Center
3100-0634-C-L	Mftg. Bldg., Old Area, Floor, South
3100-0635-C-L	Mftg. Bldg., Packing Area, Floor at Main Door
3100-0636-C-L	Mftg. Bldg., Packing Area, Floor at Packing Chute
3100-0639-C-L	Mftg. Bldg., Packing Area, Low on East Wall
3100-0640-C-L	Mftg. Bldg., Packing Area, 30 to 60 inches on West Wall
3100-0641-C-L	Mftg. Bldg., New Addition, Southwest Wall, Interior
3100-0652-C-L	Mftg. Bldg., First Floor, Southwest Floor Under Vessel
3100-0653-C-L	Mftg. Bldg., First Floor, New Addition, Center Floor by Pump

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TABLE 4.2.3.2-4
(Continued)

SAMPLE NUMBER	DESCRIPTION
<u>WIPE SAMPLES</u>	
3200-0654-C-L	Mftg. Bldg., Second Floor, New Addition, North Wall by Door
3501-0690-C-L	Mftg. Bldg., Exterior, North Wall, 0 to 24 inches by Main Door
3501-0691-C-L	Mftg. Bldg., Exterior, North Wall, 36 to 60 inches by Main Door
3502-0696-C-L	Mftg. Bldg., Exterior, South, Under Load-Out Door
3502-0697-C-L	Mftg. Bldg., Exterior, South Wall, 0 to 24 inches, Package Area Door
3502-0698-C-L	Mftg. Bldg., Exterior, South Wall, 36 to 60 inches, Package Area Door
3506-0692-C-L	Mftg. Bldg., Exterior, West Wall, 0 to 24 inches, by Large North Door
3506-0693-C-L	Mftg. Bldg., Exterior, West Wall, 36 to 60 inches, by Large North Door
3506-0694-C-L	Mftg. Bldg., Exterior, West Wall, 0 to 24 inches, by South Stairway
3506-0695-C-L	Mftg. Bldg., Exterior, West Wall, 36 to 60 inches, by South Stairway

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TABLE 4.2.5-1
COORDINATES AND ELEVATIONS OF
NEAR SURFACE SOIL SAMPLES, BORINGS, AND MONITORING WELLS

IDENTIFICATION	COORDINATES ⁽¹⁾		GROUND SURFACE
	NORTH	EAST	ELEVATION ⁽²⁾
<u>NEAR SURFACE SOIL SAMPLES</u>			
A-2-G	21.0	71.0	99.3
A-4-F	28.0	239.0	100.5
A-5-G	28.1	233.9	99.1
B-2-M	96.7	57.4	98.5
C-6-B	138.9	283.3	98.8
D-4-M	188.7	140.1	103.0
E-1-G	259.4	24.1	98.3
E-5-D	195.4	226.6	98.3
G-3-I	346.0	92.5	99.7
G-3-L	309.6	133.1	99.6
G-4-A	356.3	187.8	100.0
G-5-F	323.3	205.4	99.5
H-1-H	381.7	30.0	98.6
H-2-B	395.2	85.5	100.4
H-2-H	365.3	63.7	99.5
H-5-F	395.2	204.8	99.5
H-7-F	415.0	344.0	98.4
H-7-H	357.3	343.3	98.9
J-6-K	394.0	260.0	102.3
<u>MONITORING WELLS</u>			
MW-1A	418.2	50.0	98.7
MW-2A	440.3	214.2	98.9
MW-3A	450.4	336.9	97.3
MW-4A	147.3	339.7	97.6
MW-5A	37.7	44.6	98.9
MW-6A	14.2	137.9	98.9
MW-7A	178.0	30.3	98.4
MW-8A	280.8	335.7	99.7
MP-9A (3)	337.6	224.5	99.5

See footnotes at end of table.

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TABLE 4.2.5-1
(Continued)

IDENTIFICATION	COORDINATES ⁽¹⁾		GROUND SURFACE ELEVATION ⁽²⁾
	NORTH	EAST	
<u>BORINGS</u>			
B-1	416.7	50.6	98.7
B-2	439.3	217.3	98.9
B-3	450.0	332.3	97.3
B-4	142.8	340.2	97.6
B-5	38.3	49.7	98.9
B-6	14.1	141.6	98.9
B-7	177.4	24.9	98.4
B-8	274.3	336.1	97.7
B-9	270.2	212.2	99.3
B-10 ⁽⁴⁾	360.4	225.4	99.7
B-11 ⁽⁴⁾	313.0	228.2	99.2
B-12	348.9	224.2	99.7
B-13	346.0	224.0	98.0
B-14	-260.0	136.0	99.7
<u>EXISTING BORINGS</u>			
B-1-60	217.0	170.0	103.0
B-2-60	396.0	174.0	100.4
B-3-60	419.0	116.0	98.8
B-4-60	104.0	166.0	98.8
B-5-60	47.0	192.0	100.5

(1) Coordinates are with respect to site grid (Figure 4.2.5-1).

(2) Ground surface elevations are with respect to site datum.

(3) MP-9A is a piezometer installed in the glaciofluvial sands.

(4) Borings B-10 and B-11 were also designated as Near-Surface Soil Sample Locations G-5-E and F-5-E, respectively.

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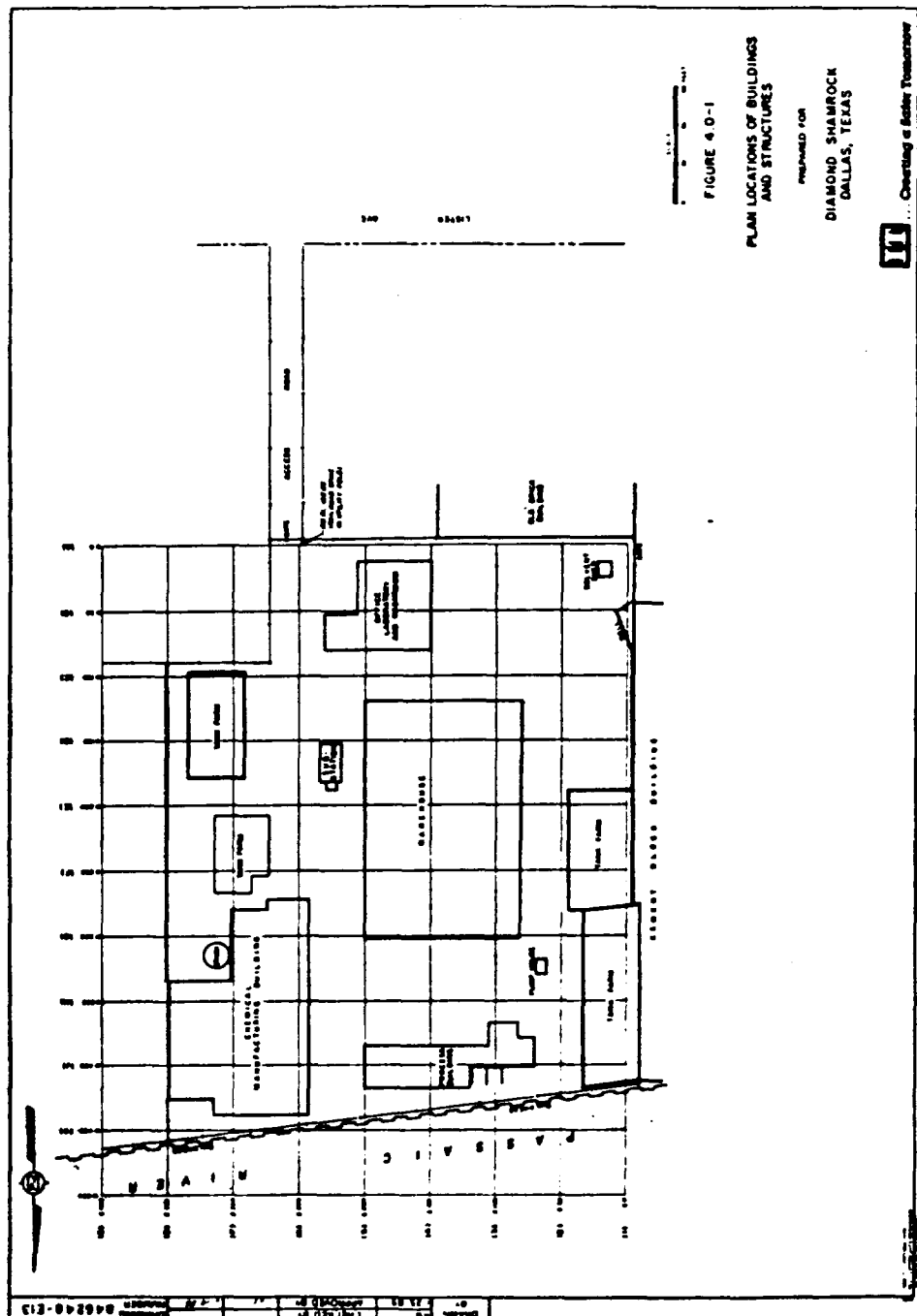
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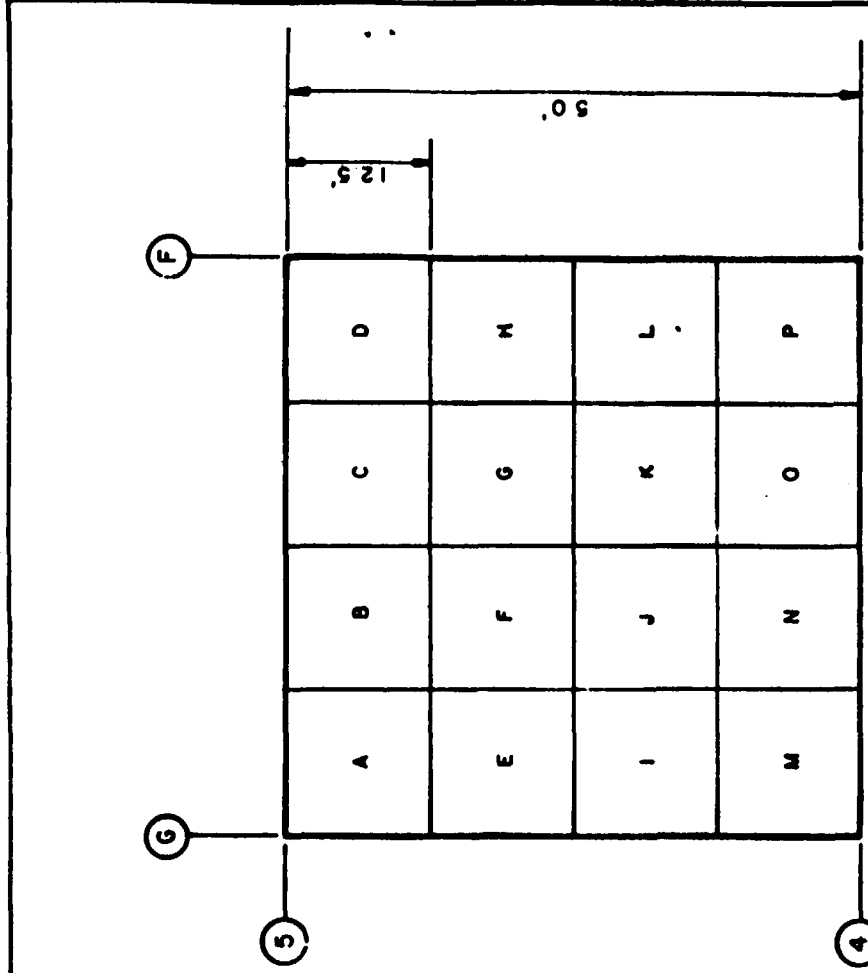


FIGURE 4.0-2

GRID SUBDIVISION
IDENTIFICATION

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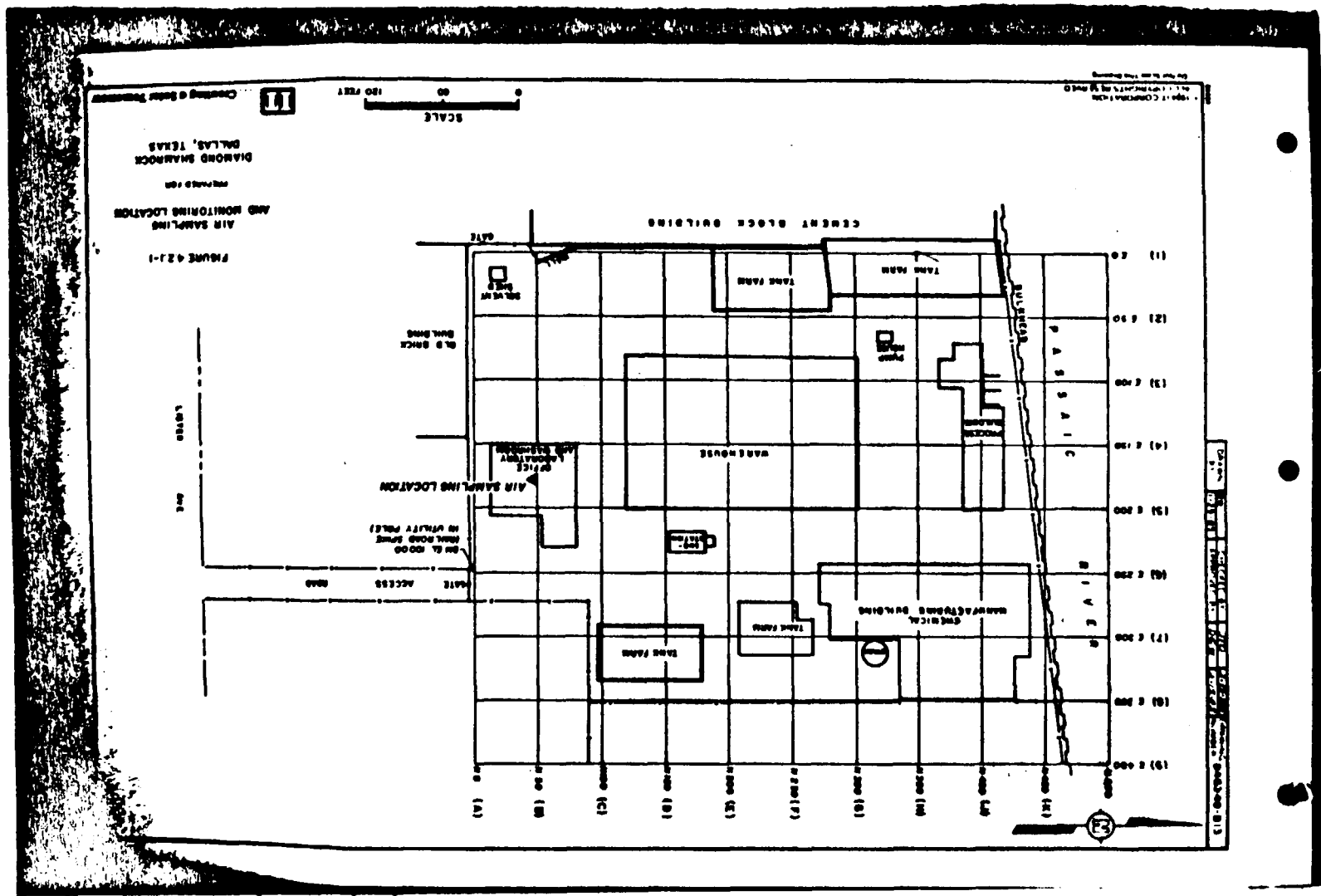


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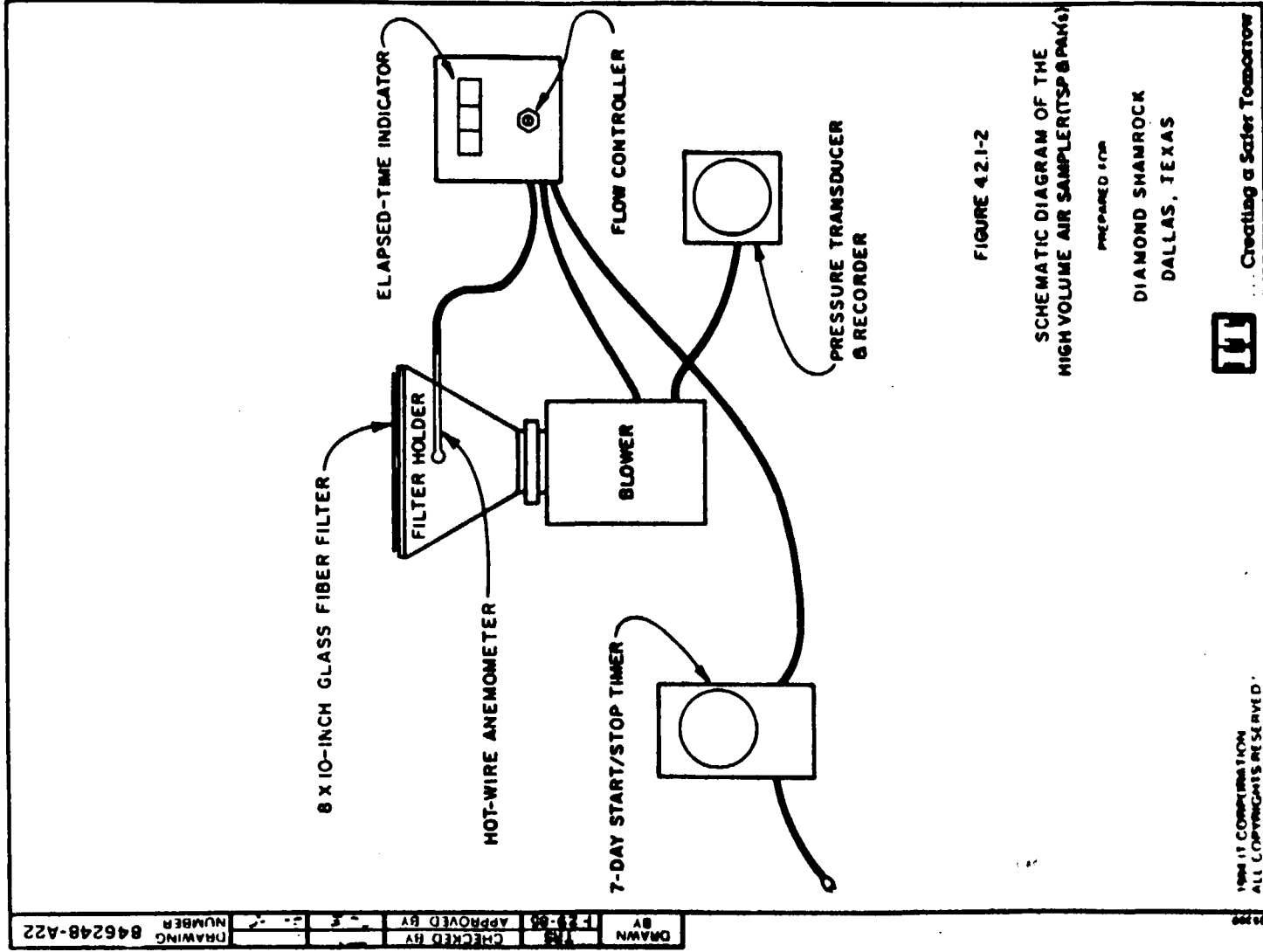


FIGURE 4.2.1-2

SCHEMATIC DIAGRAM OF THE
HIGH VOLUME AIR SAMPLER (TS-800)

PREPARED BY

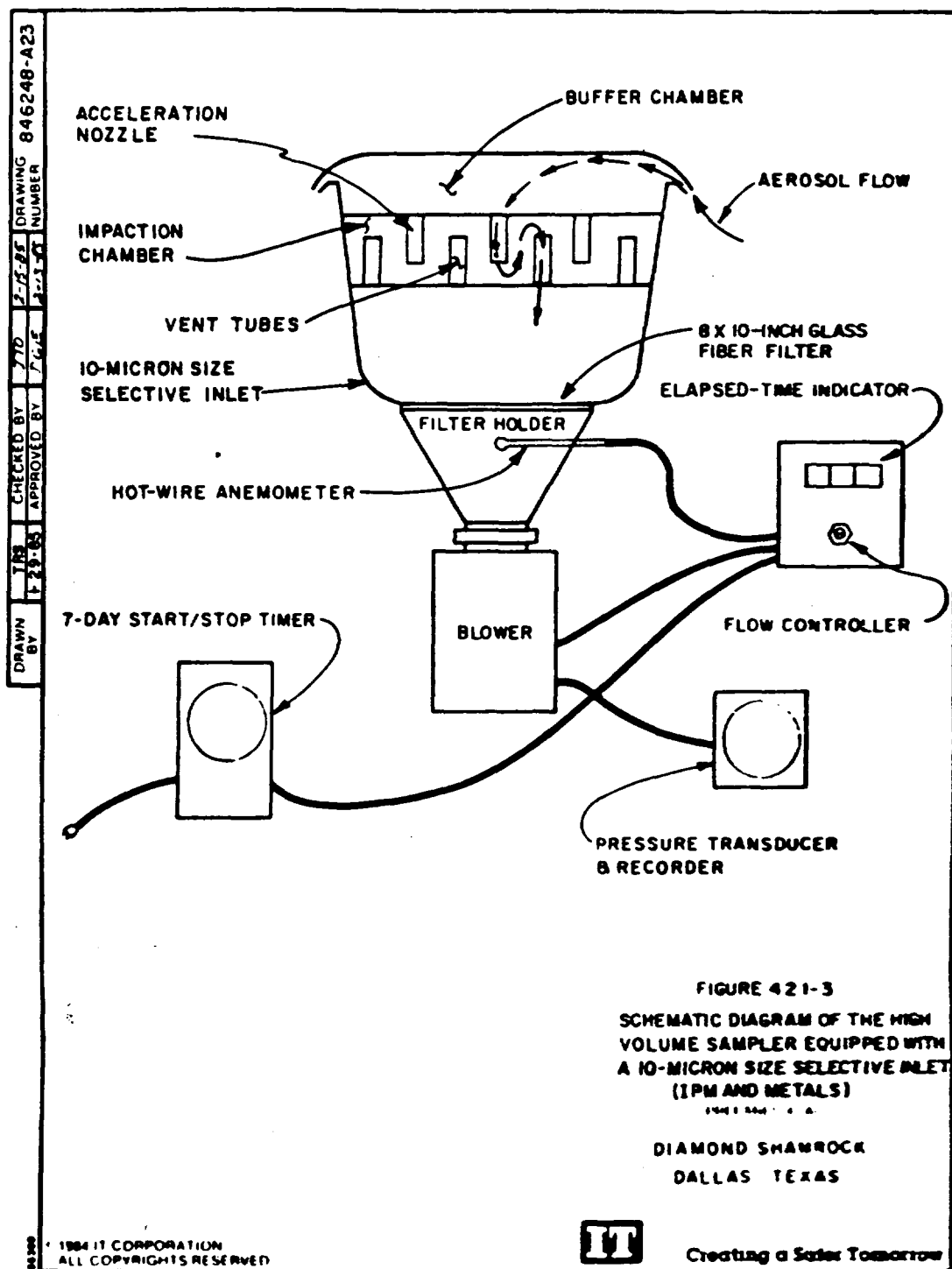
DIAMOND SHAMROCK
DALLAS, TEXAS



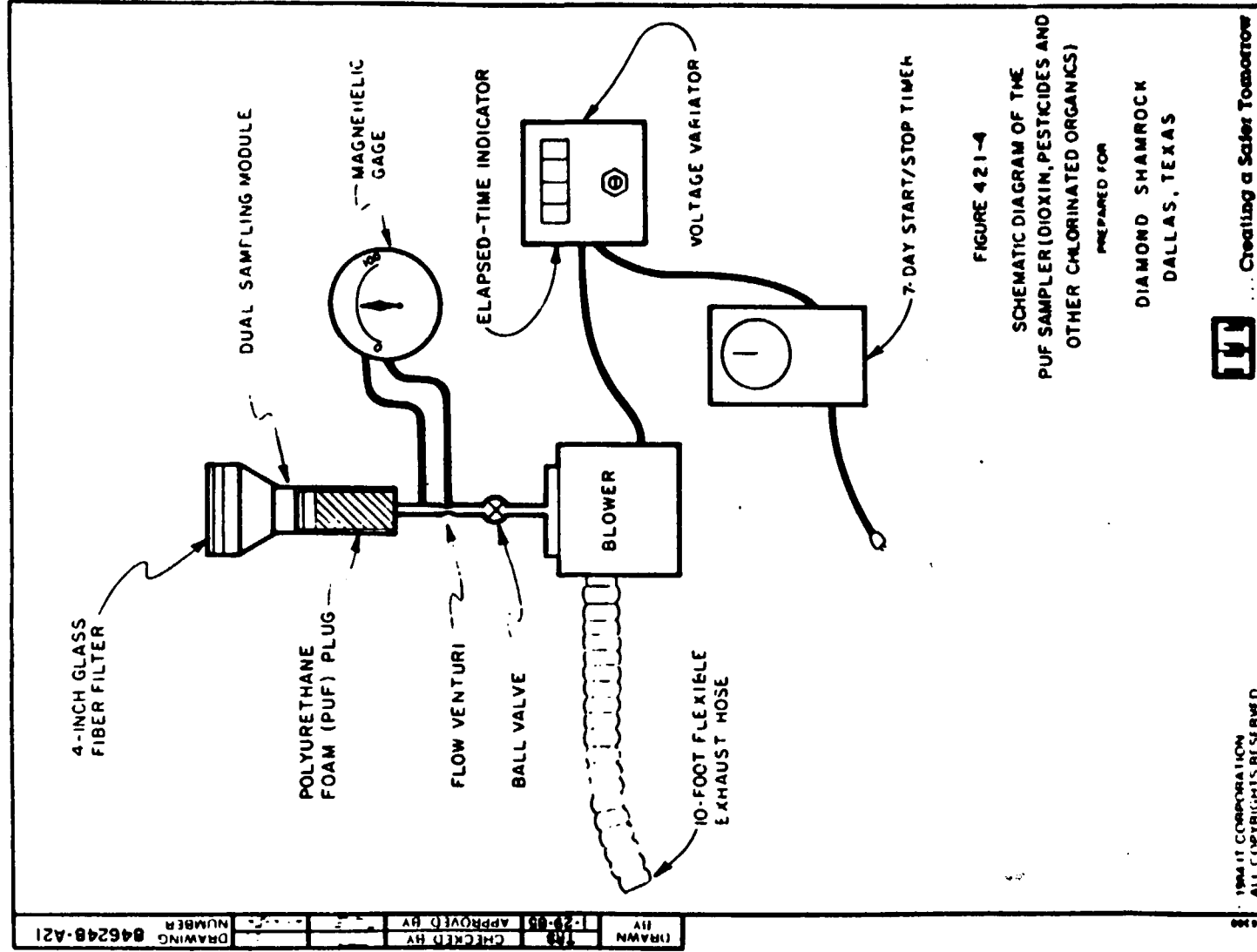
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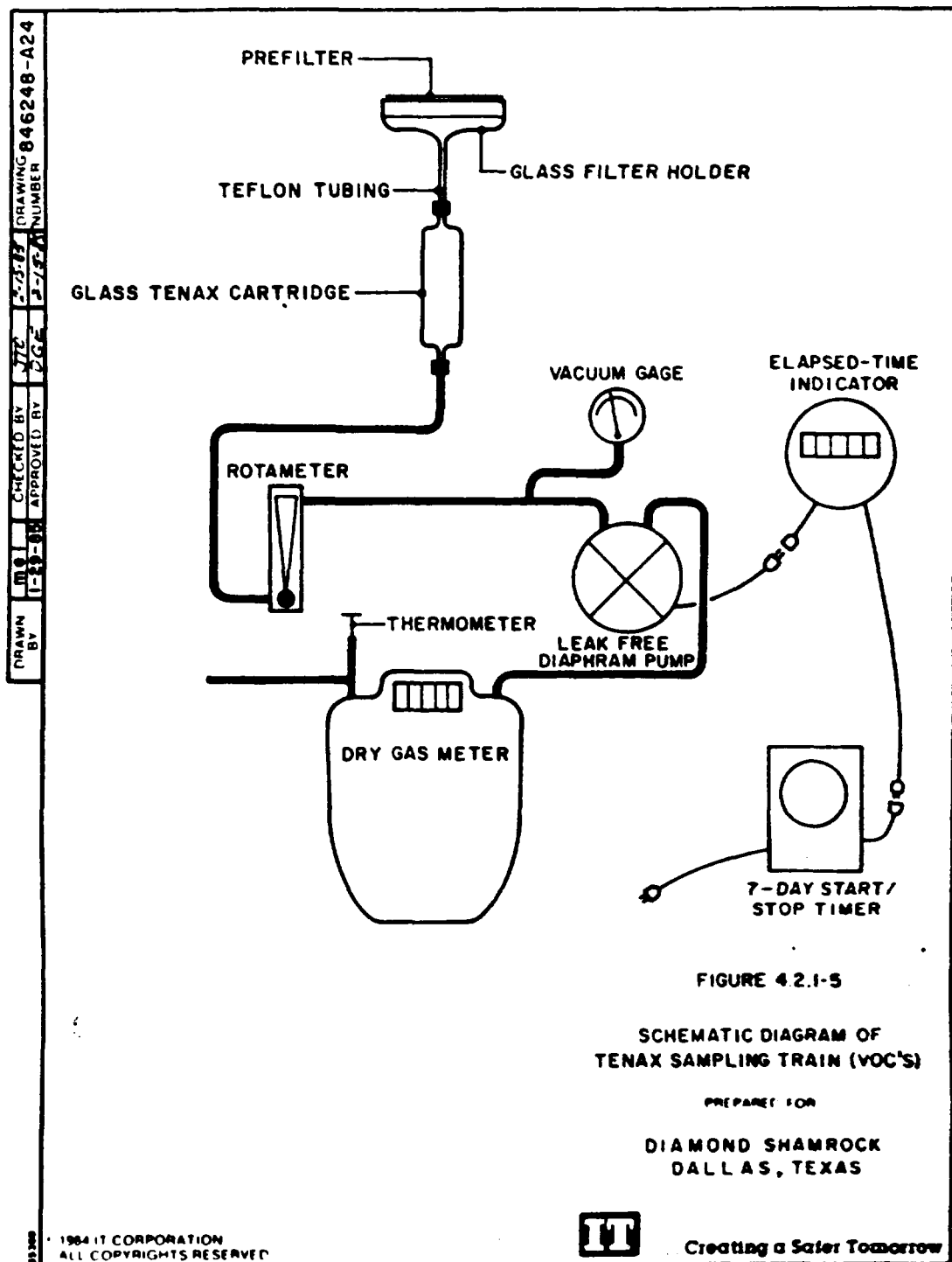
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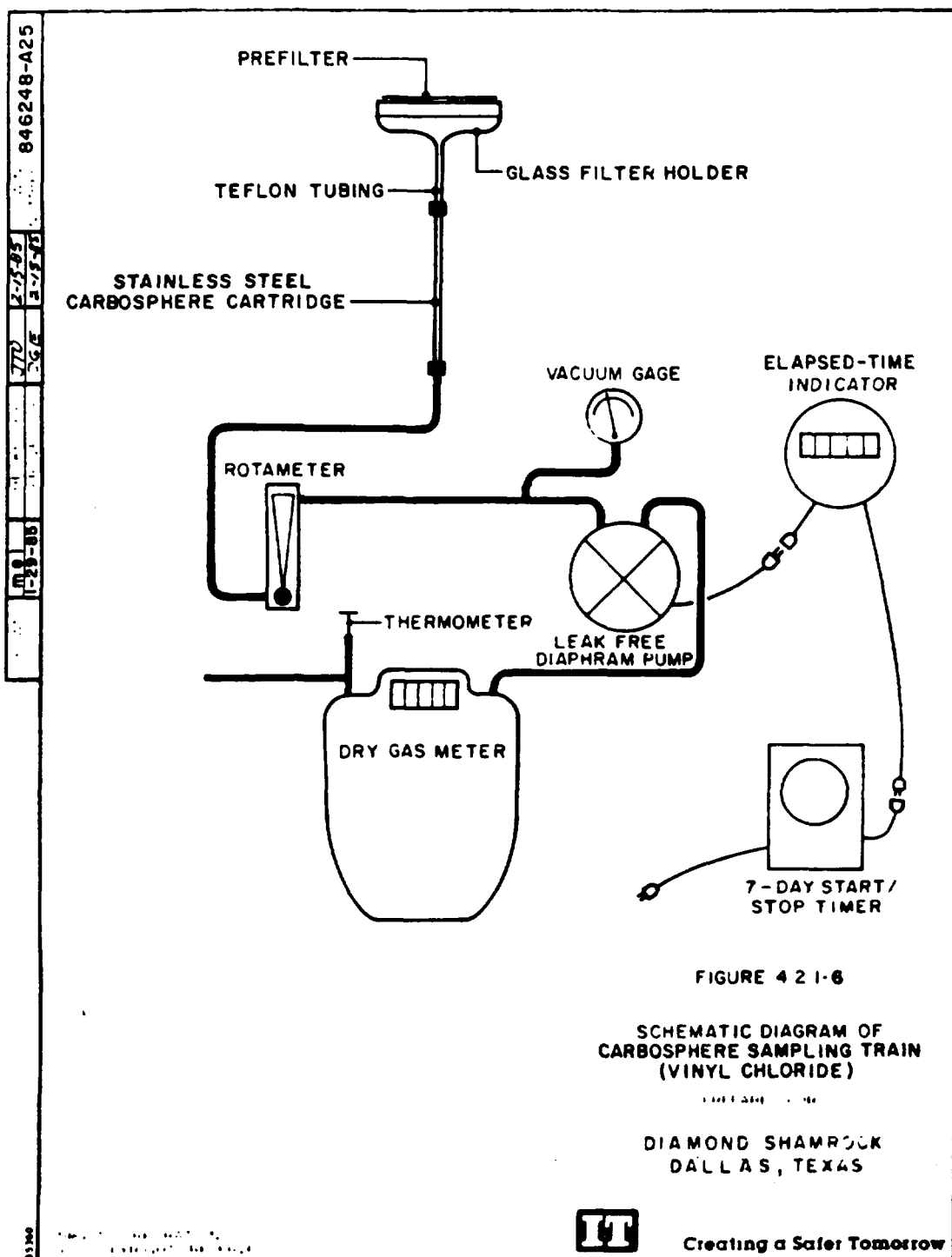


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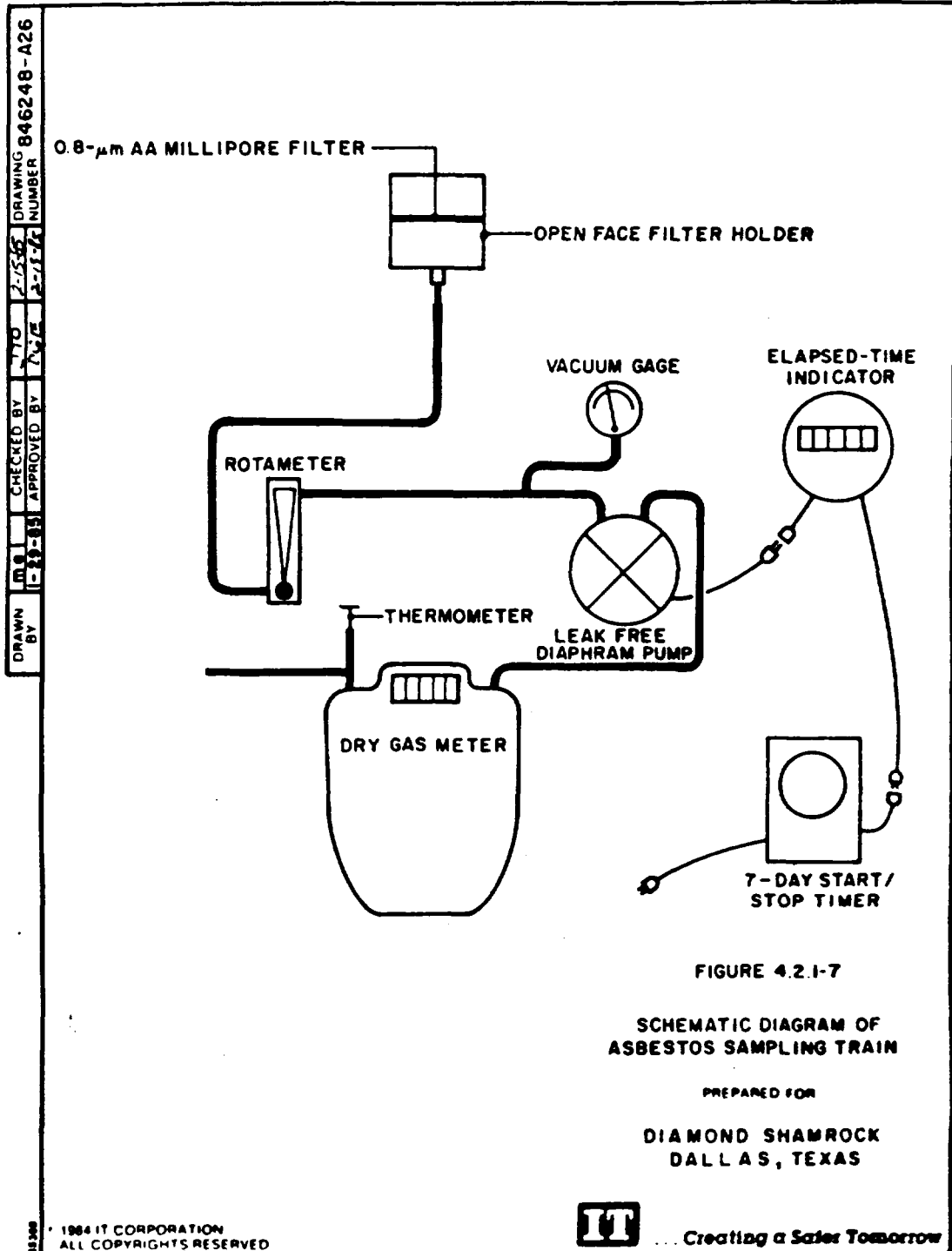
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415 MODULINE DATA SHEET

POP & POP Sample	SW & Sample Sample	POP Sample	Substrate Sample	WEC Sample	WEC	WEC
Sampler 1.D	Sampler 1.D.	Sampler 1.D.	Sampler 1.D.	Sampler 1.D.	Sampler 1.D.	Sampler 1.D.
Sample No.	Sample No.	Sample No.	Sample No.	Tube Type	Tube Type	Tube Type
Filter No. _____ Start _____ End _____ Time _____	Filter No. _____ Start _____ End _____ Time _____	Filter No. _____ Start _____ End _____ Time _____	Container No. _____ Start _____ End _____ Time _____	Sample No. _____ Start _____ End _____ Time _____	Sample No. _____ Start _____ End _____ Time _____	Sample No. _____ Start _____ End _____ Time _____
Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____	Stop _____ Date _____ Time _____
Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____	Water _____ Plant _____ Insects _____ But at last _____
Location of Plant Base _____ Soil level _____ Plant _____ Average _____	Location of Plant Base _____ Soil level _____ Plant _____ Average _____	Plant Location of _____ Soil level _____ Plant _____ Average _____	Soil level or Shading _____ Soil level _____ Plant _____ Average _____	Soil level or Shading _____ Soil level _____ Plant _____ Average _____	Soil level or Shading _____ Soil level _____ Plant _____ Average _____	Soil level or Shading _____ Soil level _____ Plant _____ Average _____
Around Plant Base _____ Average _____	Around Plant Base _____ Average _____	Around Plant Base _____ Average _____	Vertical Shading _____ Soil level _____ Plant _____ Average _____	Vertical Shading _____ Soil level _____ Plant _____ Average _____	Vertical Shading _____ Soil level _____ Plant _____ Average _____	Vertical Shading _____ Soil level _____ Plant _____ Average _____
Wood Sample Number _____	Wood Sample Number _____	Wood Sample Number _____	By the Water Shading _____ Soil level _____ Plant _____ Average _____	By the Water Shading _____ Soil level _____ Plant _____ Average _____	By the Water Shading _____ Soil level _____ Plant _____ Average _____	By the Water Shading _____ Soil level _____ Plant _____ Average _____
Comments:			By the Water Temp _____ Soil level _____ Plant _____ Average _____	By the Water Temp _____ Soil level _____ Plant _____ Average _____	By the Water Temp _____ Soil level _____ Plant _____ Average _____	By the Water Temp _____ Soil level _____ Plant _____ Average _____
Number Samples taken and measured			Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____
POP			Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____
POP			Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____
POP			Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____	Soil level or Plant _____ Soil level _____ Plant _____ Average _____

FIGURE 42 1-8

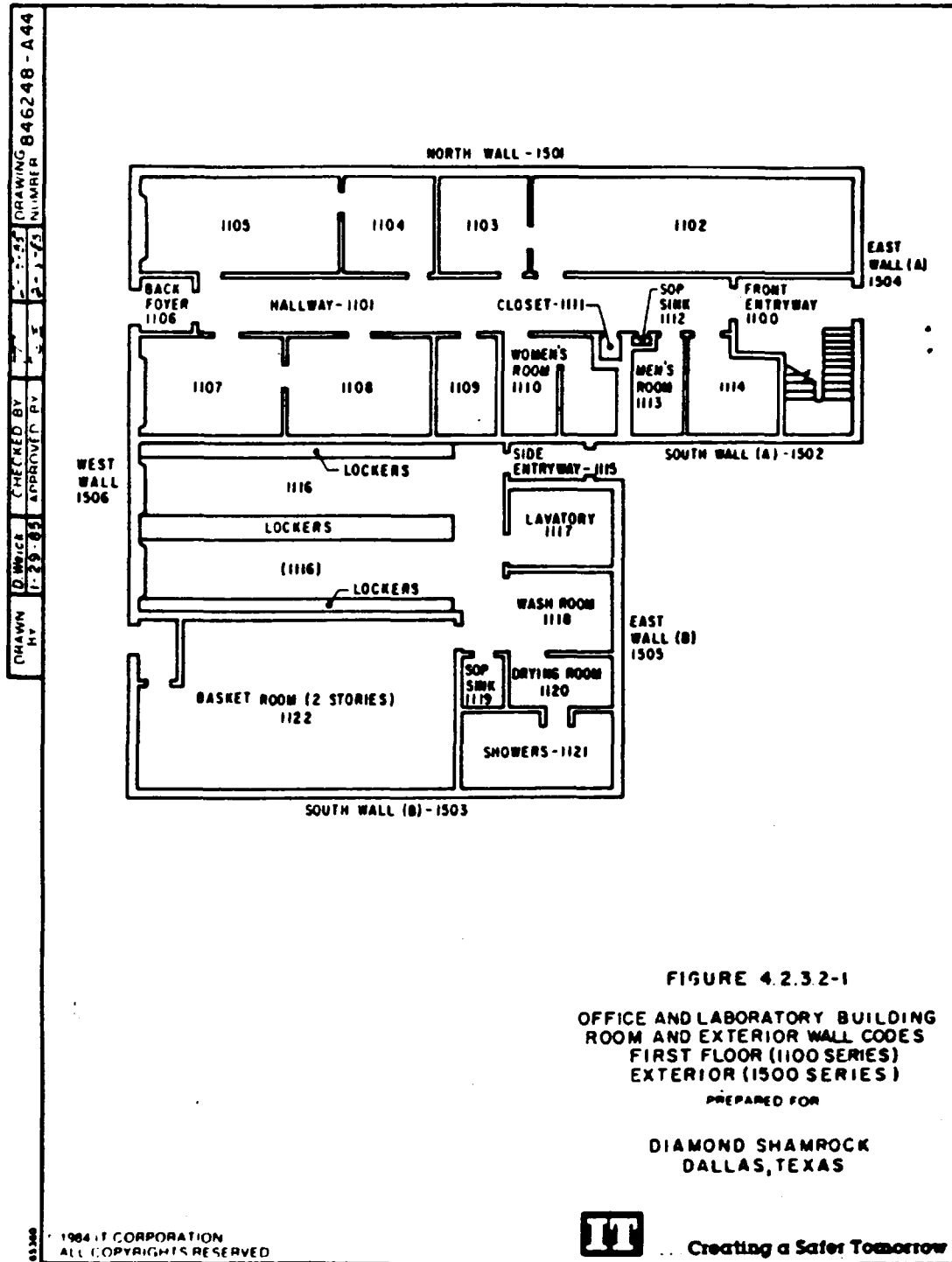
AIR MONITORING DATA SHEET

FORM PL-450 (2-7-69)

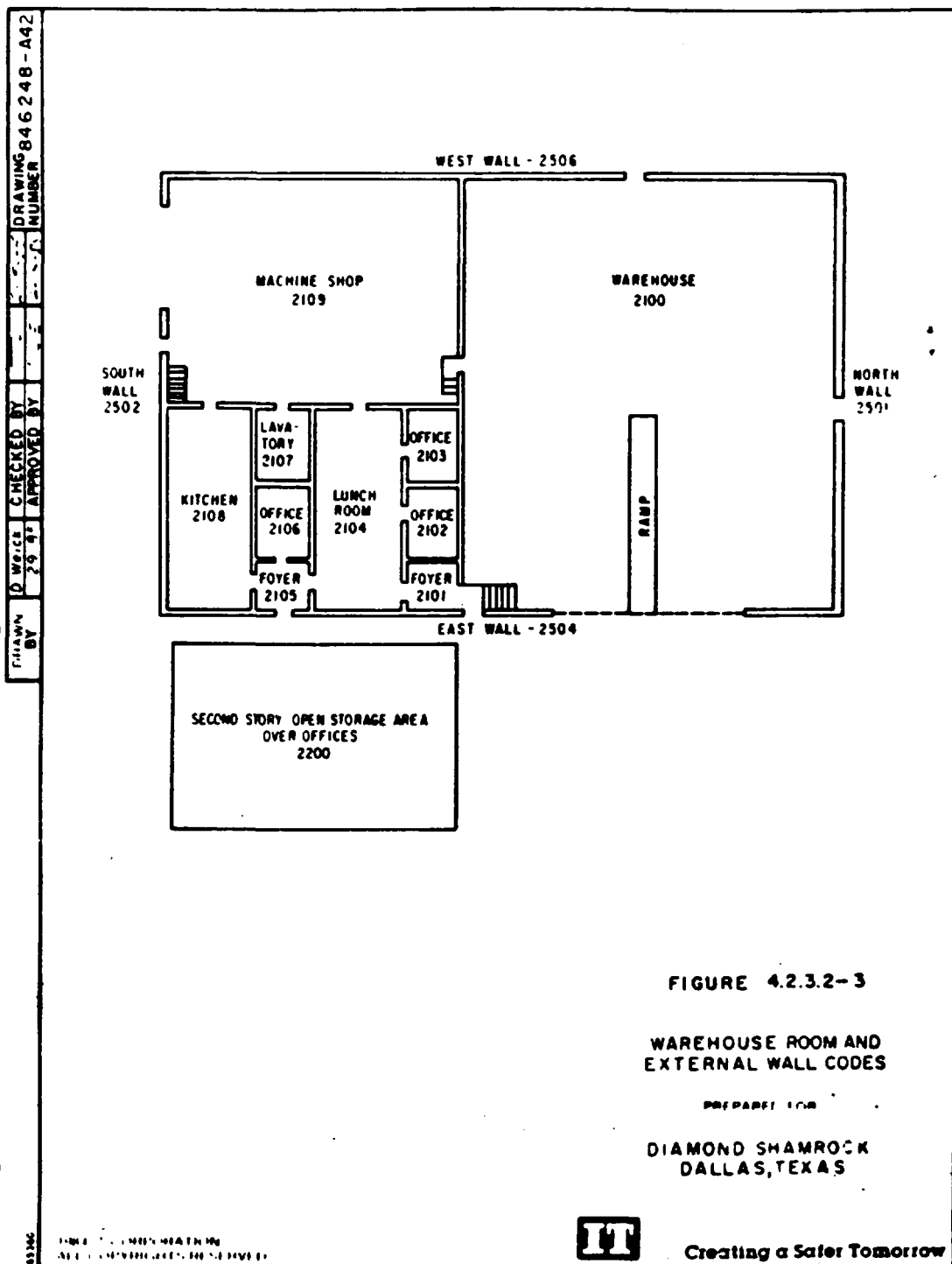
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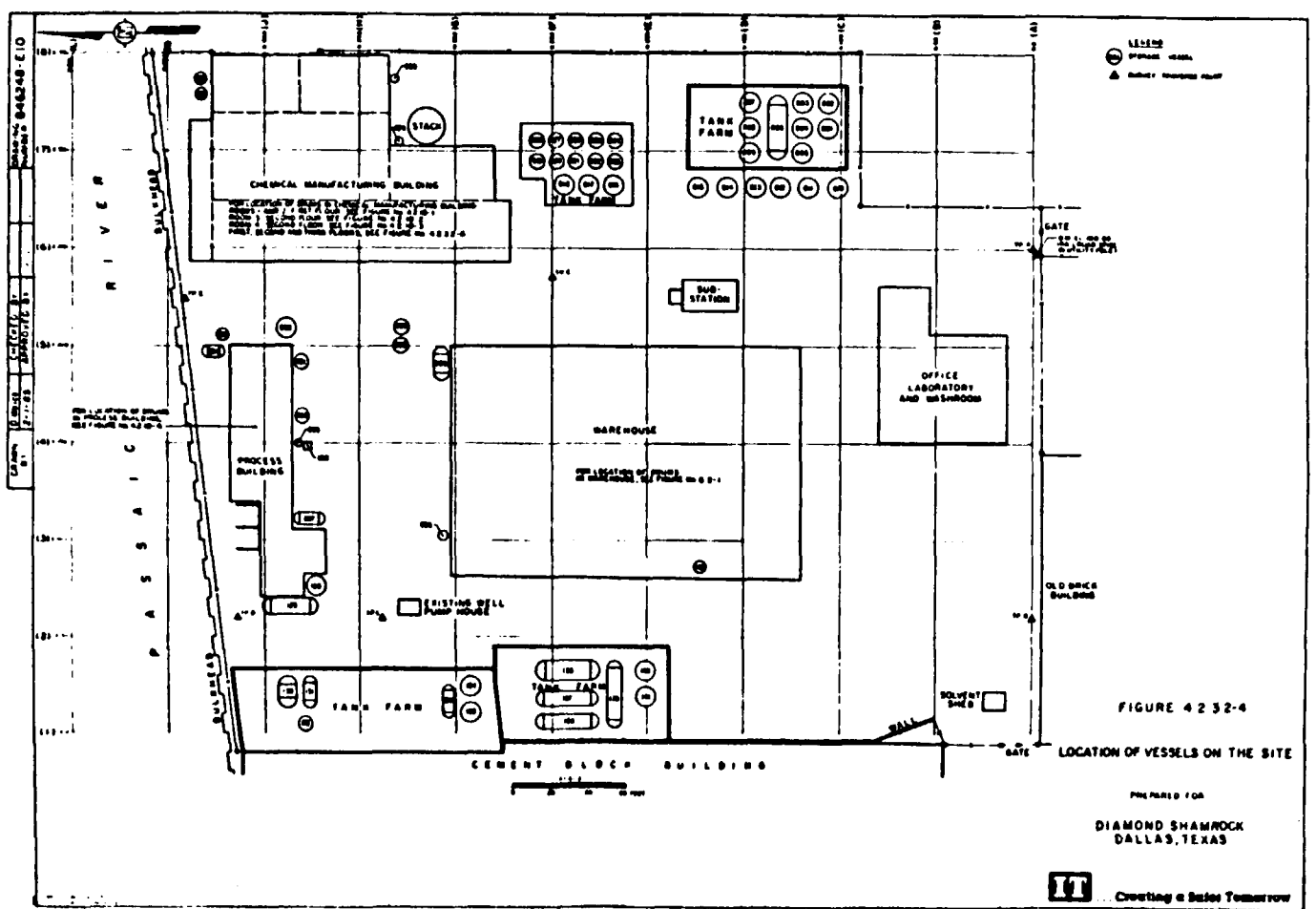
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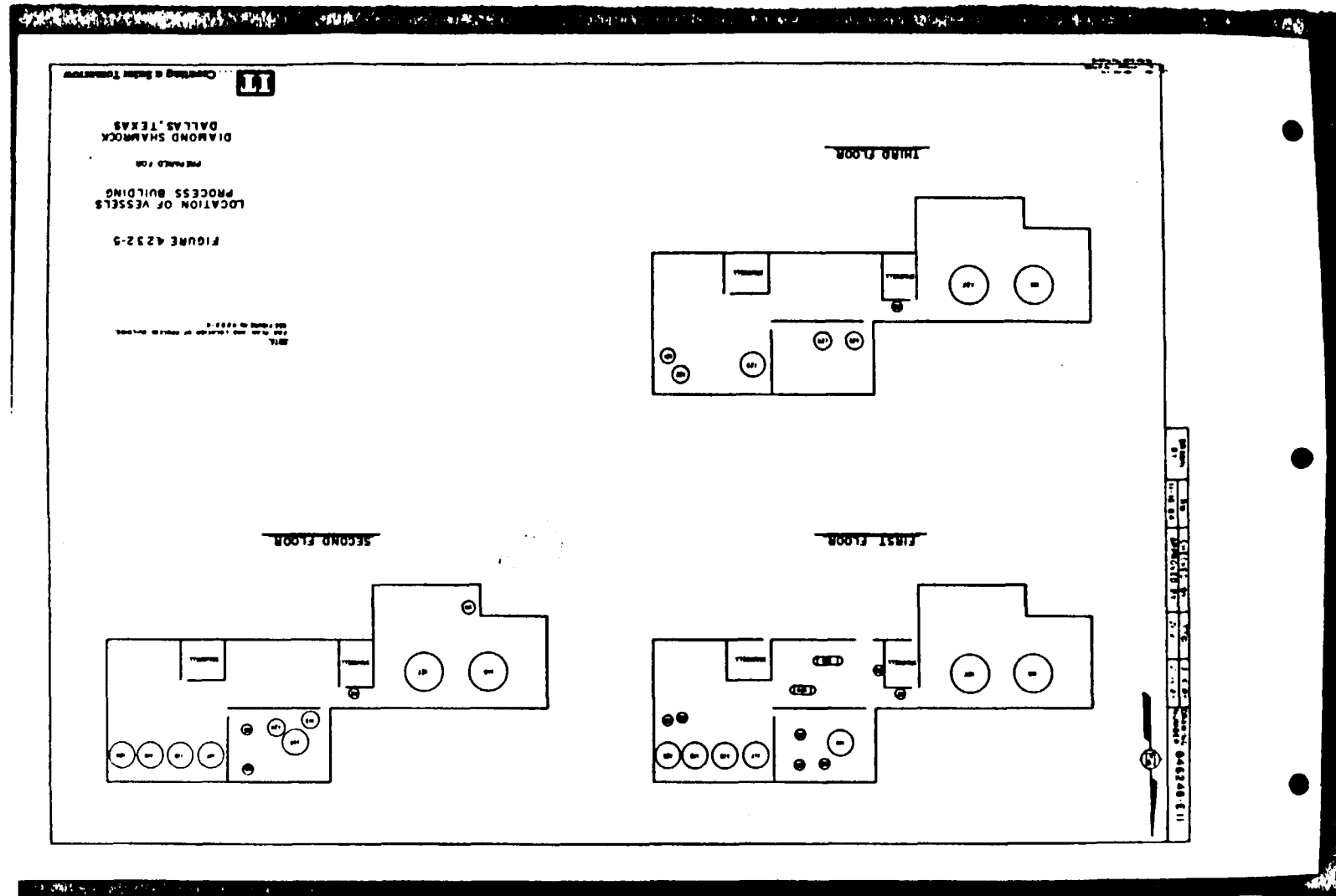
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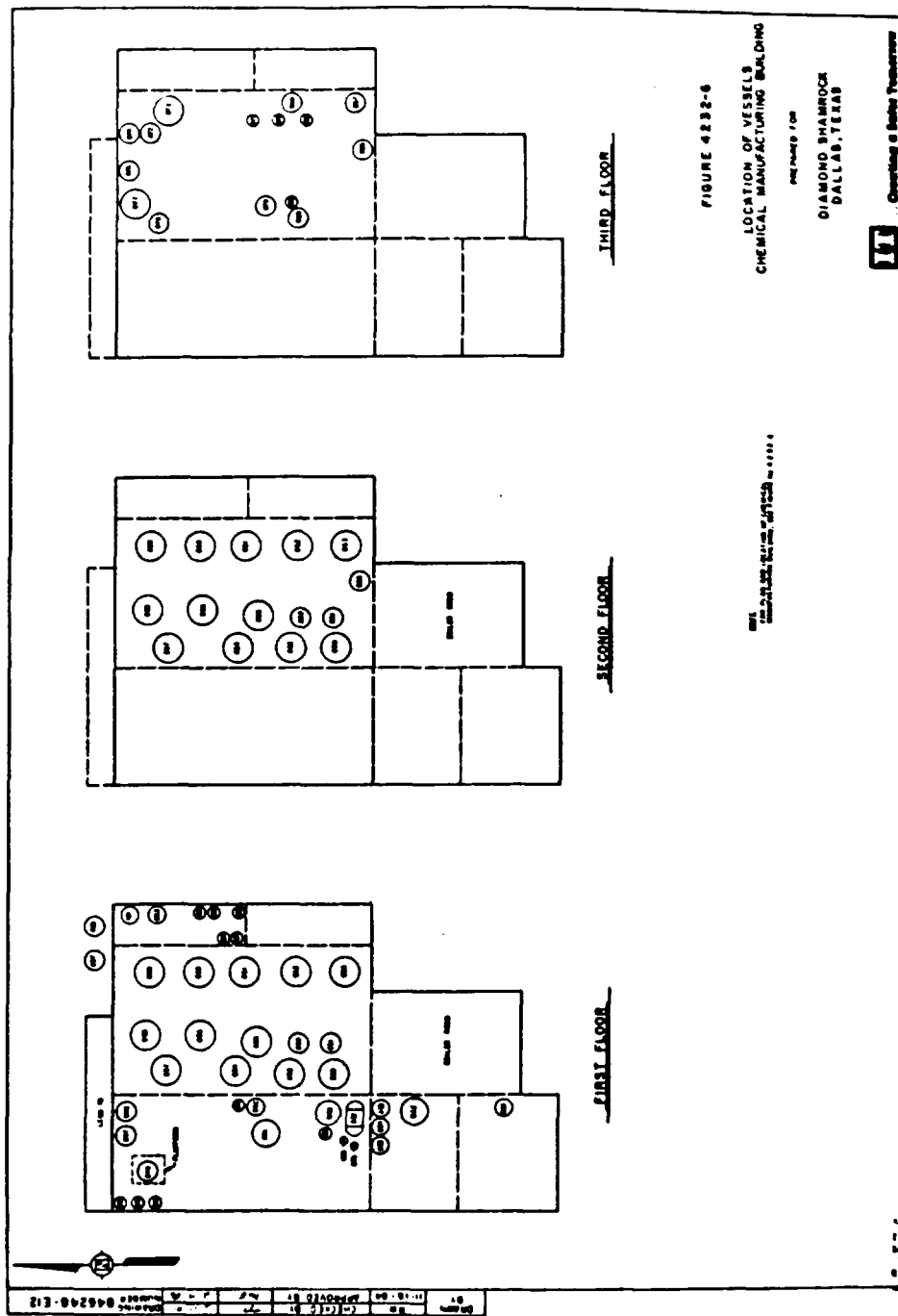
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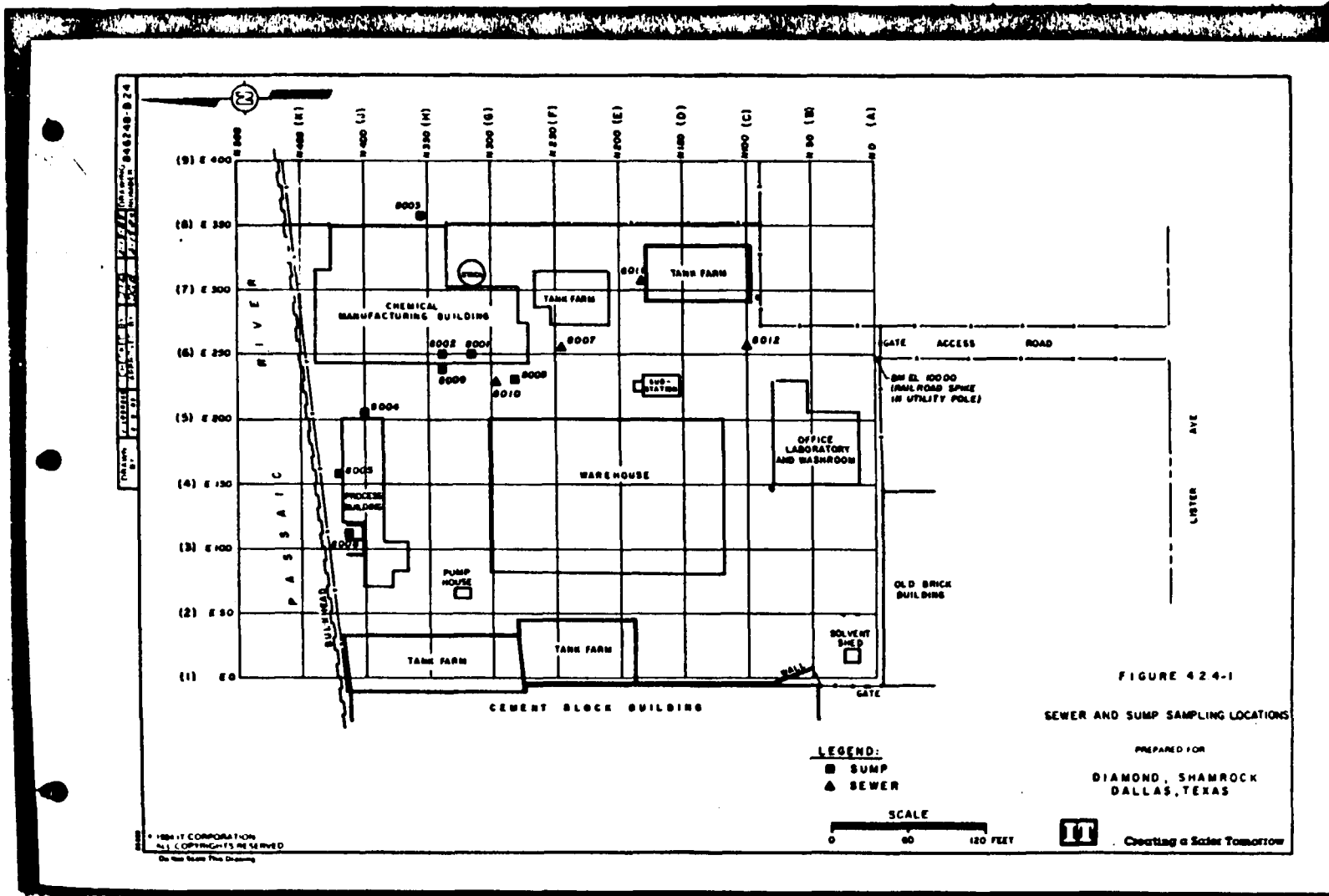


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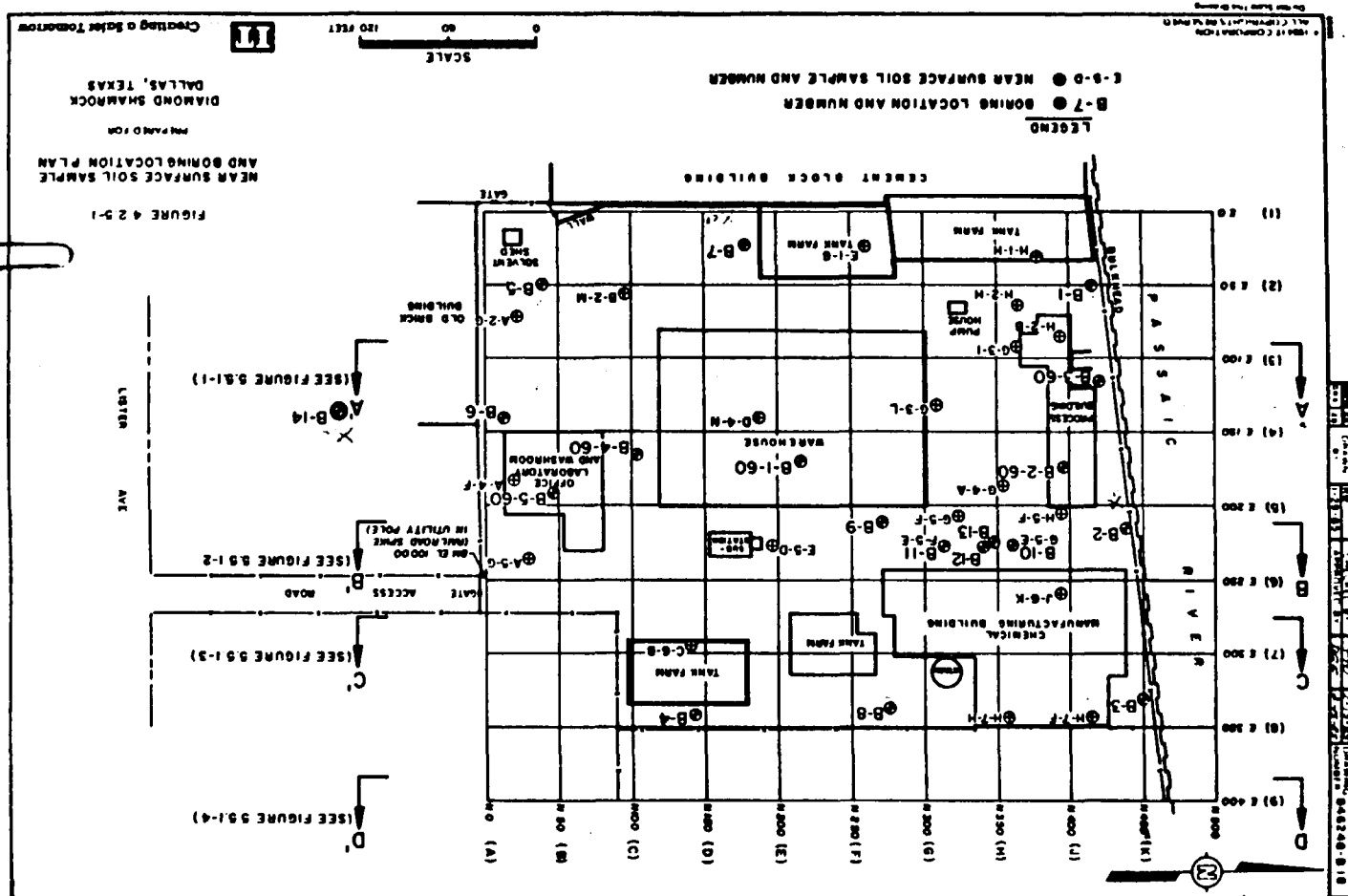
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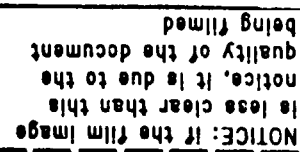
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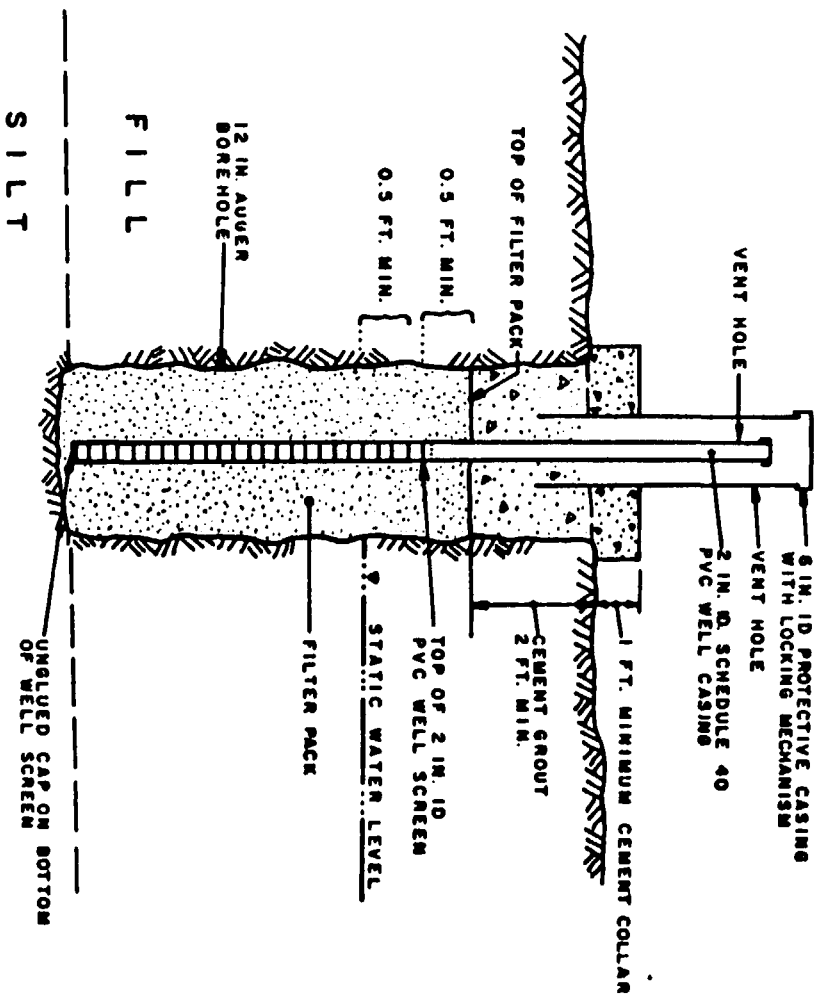


FIGURE 4.2.6.1-1

TYPICAL SHALLOW(A)
MONITORING WELL IN FILL

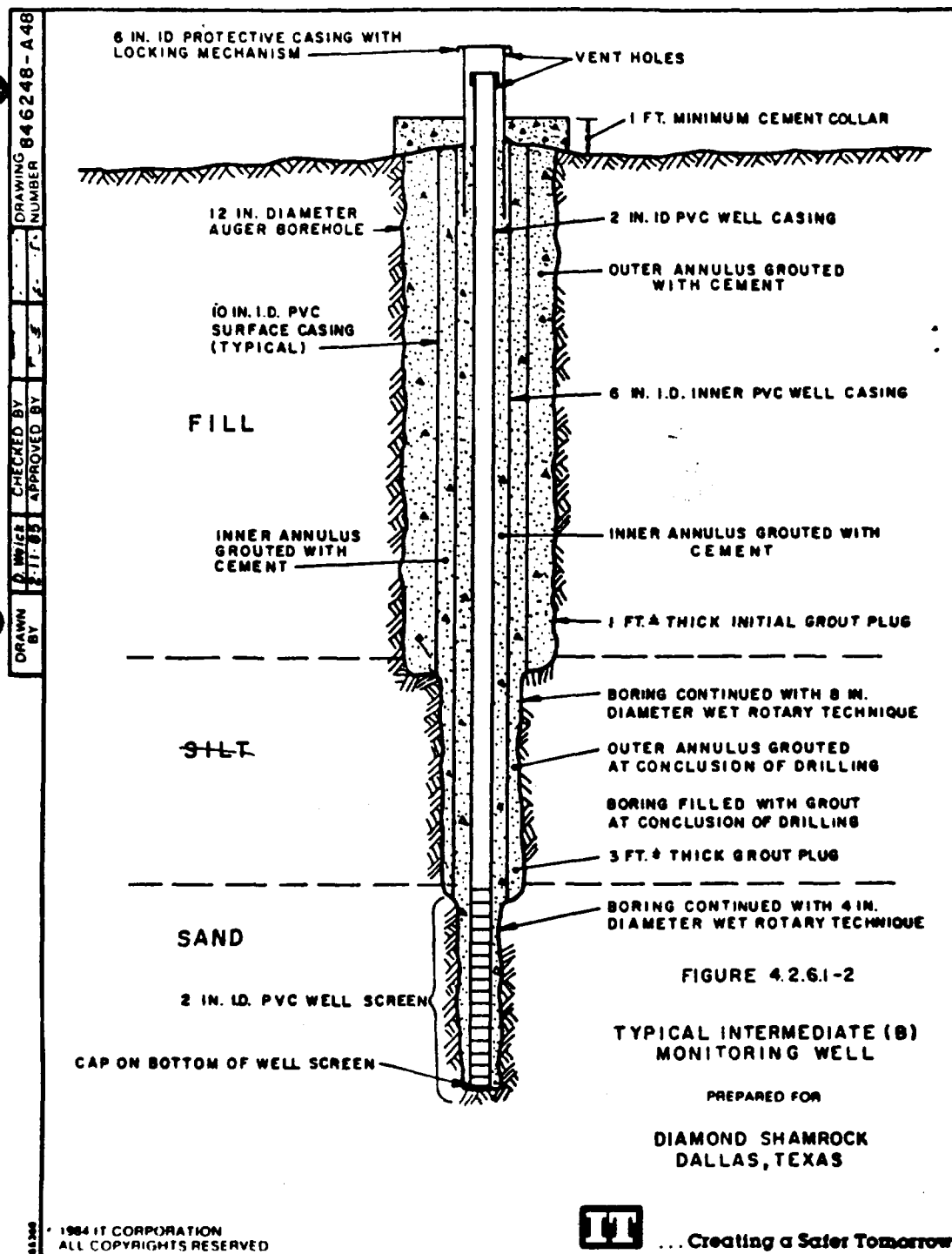
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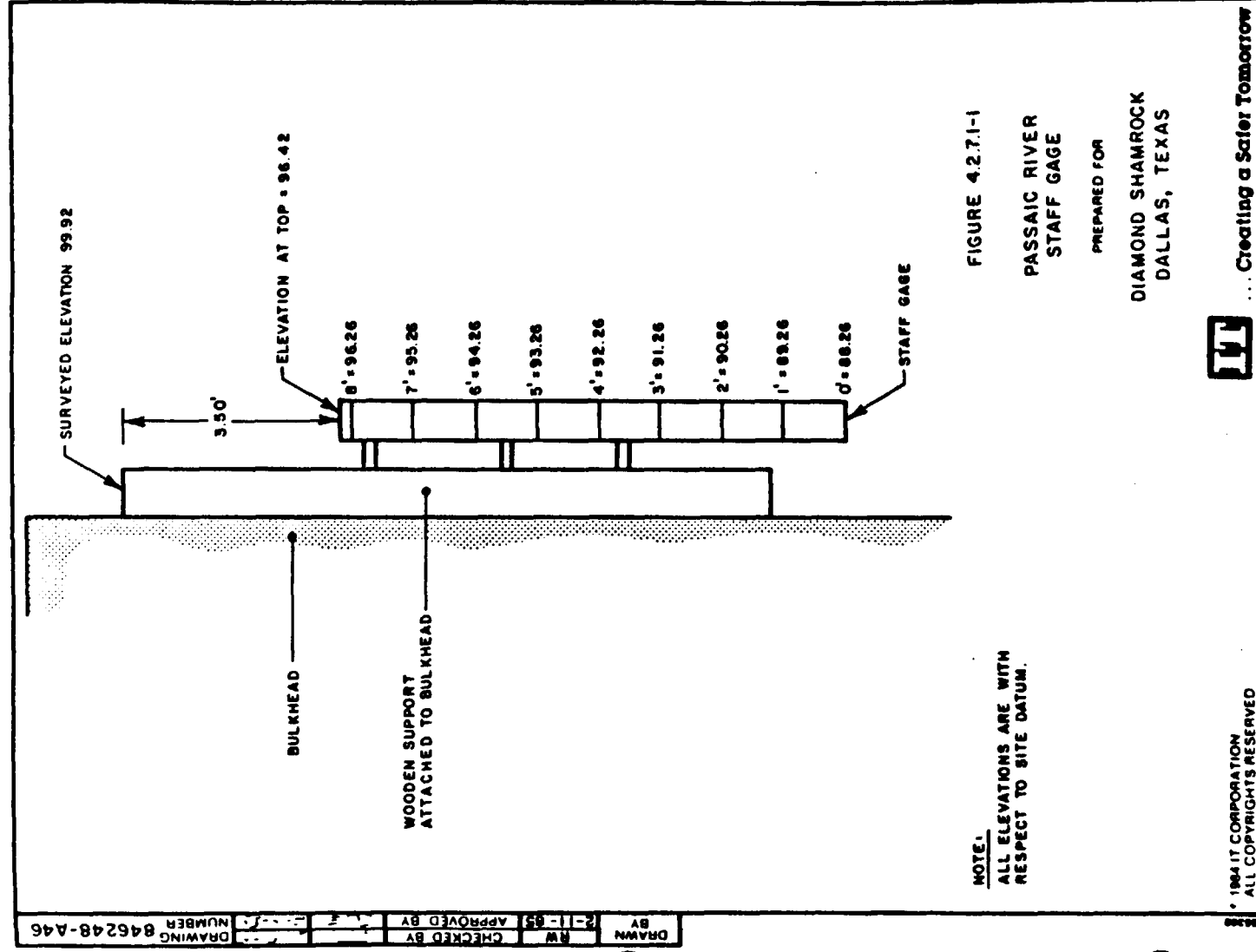
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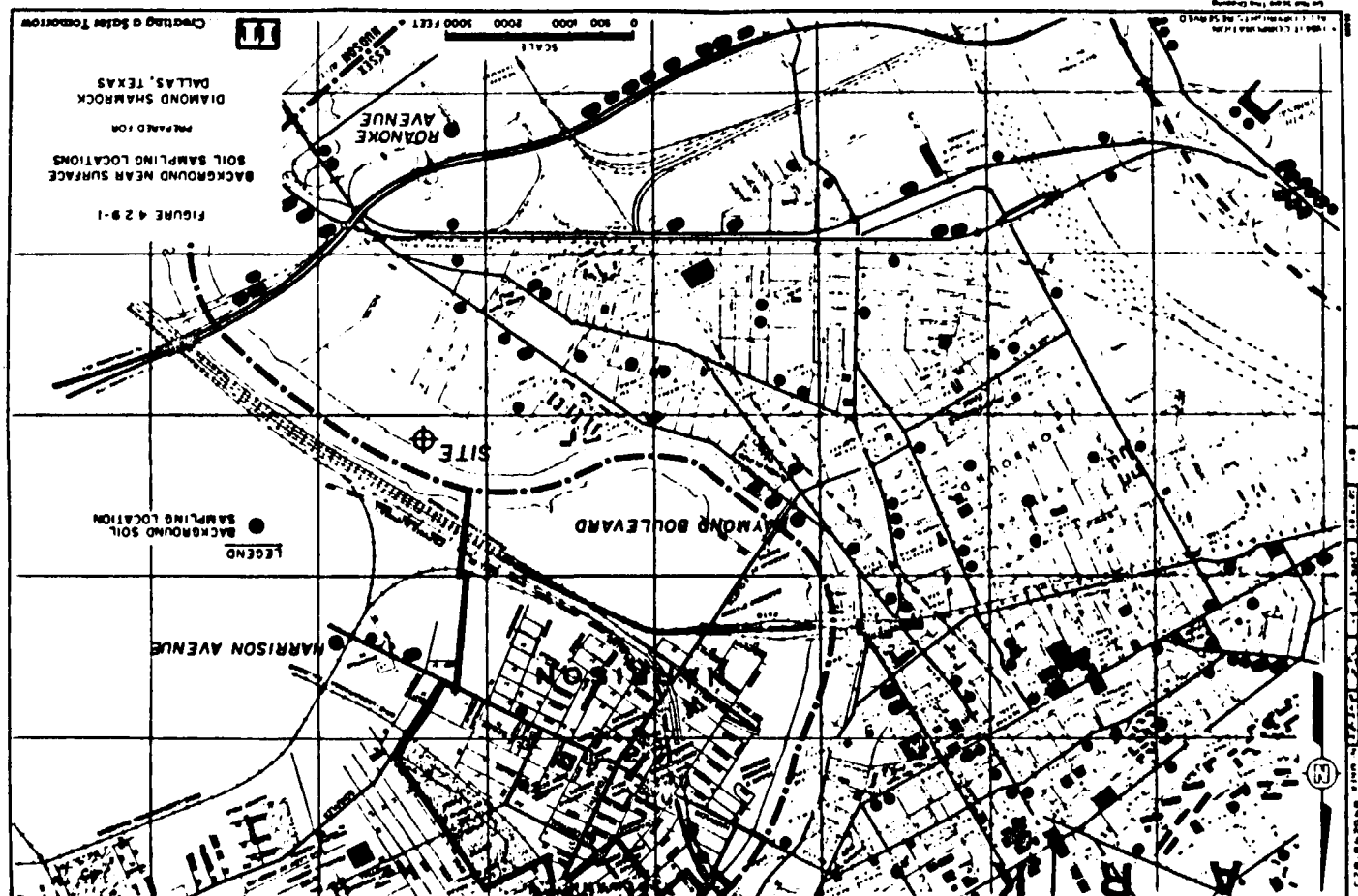
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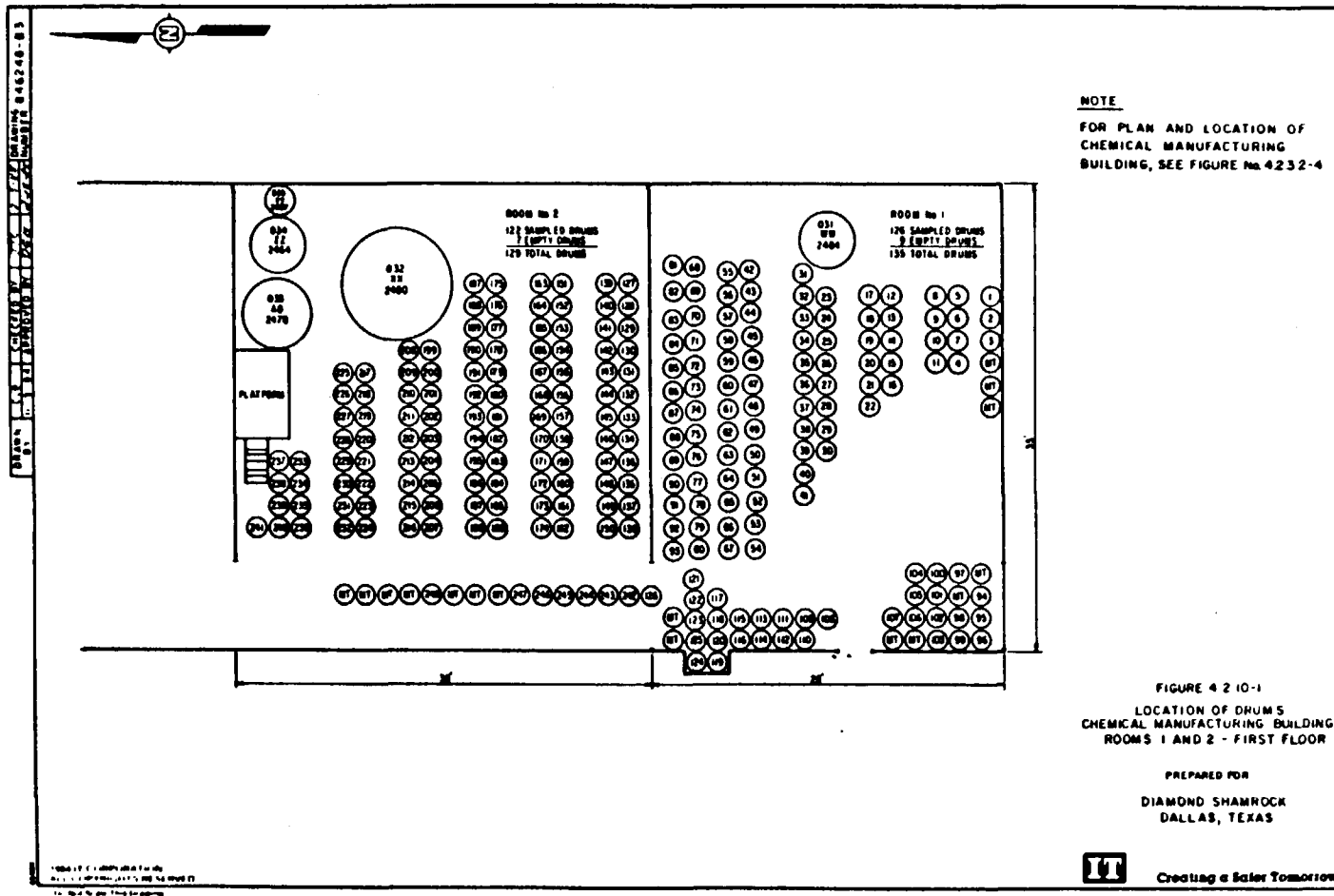
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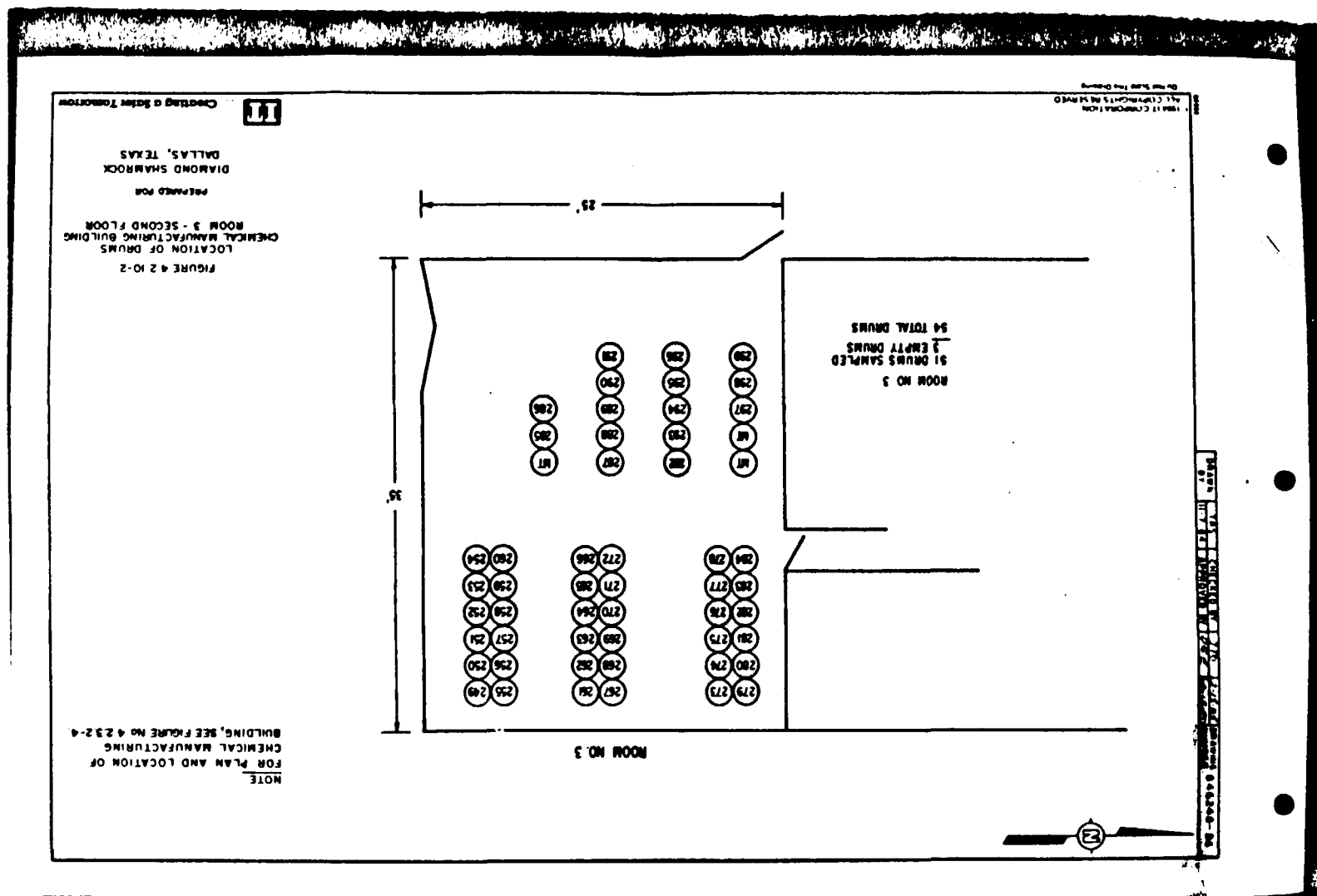
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FOR PLAN AND LOCATION OF
CHEMICAL MANUFACTURING
BUILDING, SEE FIGURE No. 4.2.32-4.

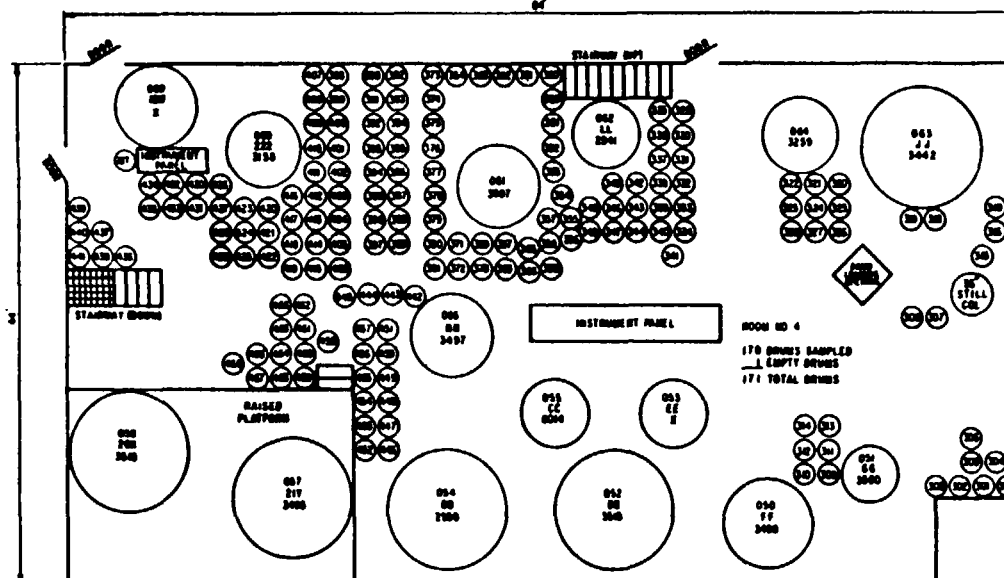


FIGURE 42 10-3

LOCATION OF DRUMS
CHEMICAL MANUFACTURING BUILDING
ROOM 4 - SECOND FLOOR

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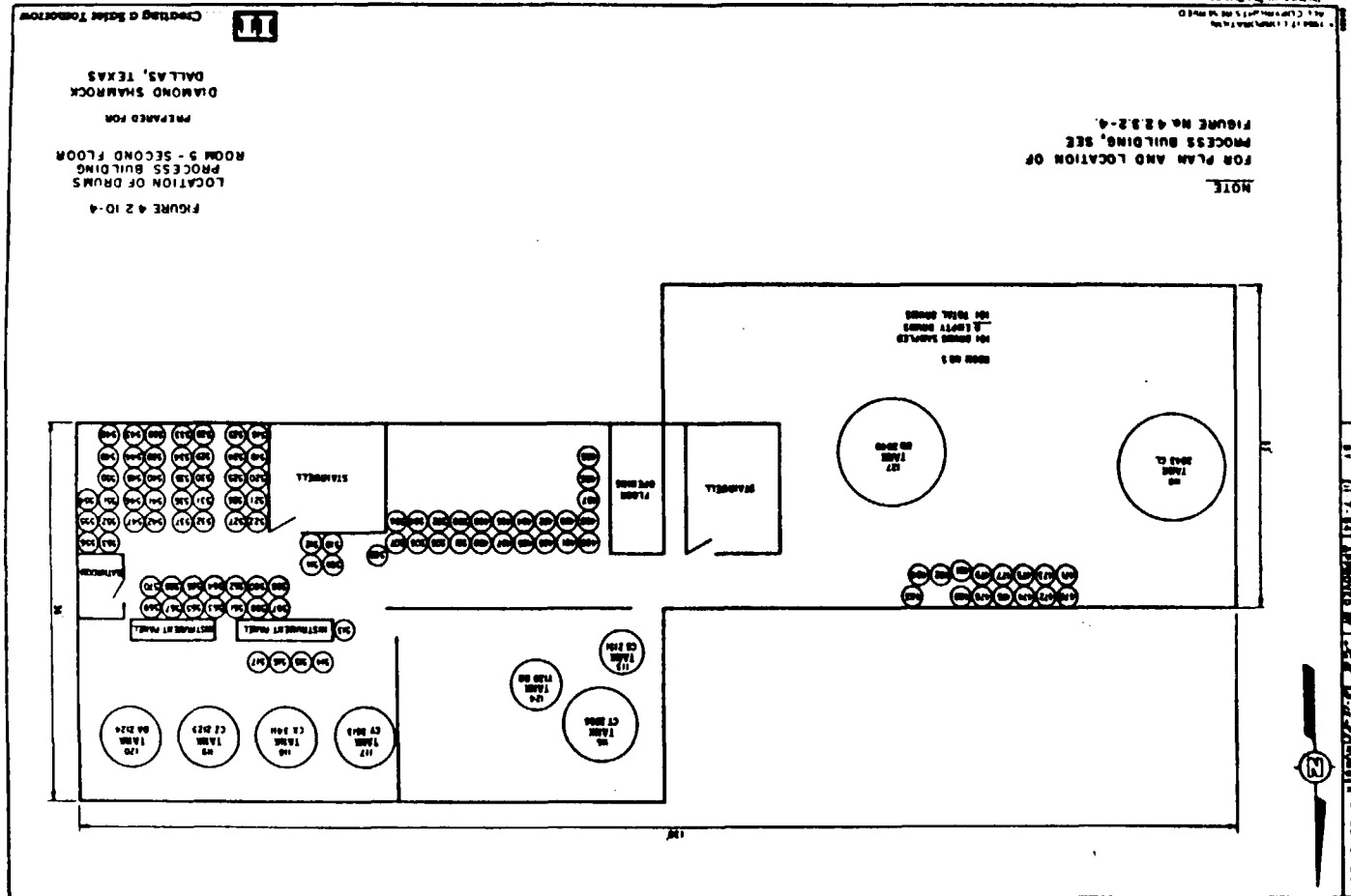
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5.0 DATA PRESENTATION

The results of the field investigation program and subsequent laboratory analytical testing are presented in the following subsections. The results include logs of borings and near-surface sample holes, hydraulic conductivities and ground water flow rates, and concentrations of priority pollutants and dioxin detected in buildings, soils, ground water and river sediments.

Of the 1540 samples collected, 877 were archived for possible future dioxin analysis. There were 532 dioxin analyses (a complete tabulation of the dioxin results is presented in Appendix C) and 122 priority pollutant analyses performed including QC, industrial hygiene and ambient air. The results of these analyses are summarized in tables and illustrated with figures presented herein. Complete laboratory analysis summaries and other supporting data are presented in appendices at the end of this report. Files of raw data as reported from the laboratories are being provided separately to NJDEP.

5.1 AMBIENT AIR

Ten sets of ambient air samples were subjected to detailed chemical analysis. As requested by the NJDEP, those sets of samples having the ten highest iron and manganese concentrations were analyzed. Table 5.1-1 presents the results of the inhalable particulate matter (IPM) filter analyses for iron and manganese (in total micrograms per filter), and indicates those sample sets chosen for detailed chemical analysis. One of the sample sets chosen corresponds to a one-in-six day sampling normally conducted by NJDEP. Five such NJDEP sampling days occurred during the site investigation. At the request of the NJDEP, the selected sample list was modified by substituting a set of samples corresponding to another NJDEP sampling period (September 21, 1984) for the originally selected October 8, 1984 set. The sample sets are identified in Table 5.1-1. The analytical results of the ambient air sampling are reported in Tables 5.1-2 through 5.1-7.

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Meteorological data were collected at the ambient air monitoring site throughout the sampling period. Data were also obtained from the weather station located at the Newark International Airport (approximately three miles southwest of the site) for the sampling period. These data include temperature, barometric pressure, sky cover, precipitation, and wind speed and direction. To date only wind direction data have been summarized. Figure 5.1-1 is a wind rose of the wind direction data collected at the site from September 7, 1984, through October 9, 1984. Figure 5.1-2 is a similar wind rose for the same period using data from National Weather Station (WSO) located at Newark Airport. Differences in the appearance of the two wind roses can be attributed to local topographic features affecting the wind flow patterns.

5.2 INDUSTRIAL HYGIENE

During the site investigation, a comprehensive industrial hygiene monitoring program was conducted so that the adequacy of the assigned worker protection could be evaluated and demonstrated. The results of the program are discussed below.

5.2.1 Atmospheric Samples for Dioxin

Table 5.2.1-1 presents the results of all dioxin related industrial hygiene monitoring; of these, 43 were atmospheric samples. For seven of the atmospheric samples, resultant concentrations were greater than the permissible exposure level (PEL) of 0.5 nanograms per cubic meter of air (ng/m^3). None of the 24 blank samples indicated detectable levels of dioxin. The seven samples are associated with four specific operations. The operations and the actions taken were:

- o Chip sampling of outside walls produced airborne concentrations of dioxin of $0.74 \text{ ng}/\text{m}^3$ and $1.23 \text{ ng}/\text{m}^3$ on September 14 and 18, 1984. All other monitoring during chip sampling resulted in concentrations below detectable levels. Because of the two results above the PEL, all future chip sampling will require the use of self-contained breathing apparatus (SCBA).

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5.2.3 Atmospheric Samples for Volatile Organics, Semi-Volatile
Organics, Alcohols, 2,4,5-T, 2,4-D, Asbestos, and Sulfuric Acid

A total of 24 samples were obtained for analysis of the above compounds. A summary of results for 2,4,5-T, 2,4-D, asbestos, sulfuric acid and all volatile and semivolatile compounds for which positive results were obtained is presented in Table 5.2.3-1. All positive results observed were of sufficiently low concentration not to be of concern.

5.2.4 Heat Stress

A Reuter Stokes Heat Stress Monitor was used on 11 days during September and October, 1984, due to the potential for heat stress related illness on these days. Results were compared to the American Conference of Governmental Industrial Hygienists recommended threshold limit value (TLV). A work-rest regimen was established at those times when the TLV was exceeded.

5.2.5 Noise Monitoring

The most significant source of noise during the site investigation was the drilling rigs. Accordingly, noise measurements were taken at the drilling operations on October 17, 1984. Measurements ranged from below 80 decibels (dBA) to 84 dBA. The permissible exposure limit is 85 dBA for hearing protection requirements and 90 dBA for engineering controls. These limits were not exceeded.

5.2.6 General Health and Safety

The procedures presented in the Work Plan were followed. All personnel entering the site were given one to two days of site specific training, depending on the amount of health and safety training previously received. All ITC employees and subcontractor employees were required to have medical examinations prior to entering the contamination zone. All personnel on site were required to wear the protective equipment prescribed in the Work Plan.

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5.3.1 Office and Laboratory Building

Figures 4.2.3.2-1 and 4.2.3.2-2 show the first- and second-story floor plans of the office and laboratory building and the room designations. Table 5.3.1-1 presents the dioxin analysis results for wipe and chip samples collected in the office and laboratory building. Of the 39 samples collected in this building, dioxin was not detected in seven and one sample was voided. Dioxin concentrations of the first floor wipe samples ranged from 38 to 1,100 ng/m². Dioxin concentrations of first floor chip samples ranged from 2.0 to 69.3 ppb. Fifteen of 15 first-floor samples were identified as containing dioxin. Dioxin concentrations of the wipe samples taken on the second floor ranged from 10 to 14,000 ng/m² with 11 of the 11 valid samples having dioxin identified. The dioxin concentrations of chip samples from the exterior of the building ranged from 0.57 to 2.4 ppb with 5 of 11 samples having dioxin identified. One exterior wipe sample had a dioxin concentration of 168 ng/m²; the other showed no dioxin present.

5.3.2 Warehouse

Figure 4.2.3.2-3 shows the floor plan of the warehouse with the assigned room designations. Table 5.3.2-1 provides the dioxin analysis results for wipe and chip samples collected in the warehouse. Of the 24 samples collected, 21 samples showed detectable levels of dioxin. The dioxin concentrations of interior wipe and chip samples ranged from 130 to 19,000 ng/m² and from 48.7 to 192 ppb, respectively. All 11 interior samples had positive dioxin results. The dioxin concentrations of exterior chip samples ranged from 1.0 to 16.5 ppb with 9 of 12 samples having positive results. The single exterior wipe sample showed dioxin present at 13 ng/m².

5.3.3 Manufacturing Building

Table 5.3.3-1 summarizes the dioxin analysis results for all wipe and chip samples collected from the manufacturing building. Positive dioxin results were obtained for 27 of the 28 samples collected. The dioxin

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Table 5.3.6-1 presents the dioxin results for the 12 tank samples analyzed, of which nine (75 percent) had positive dioxin results. Three samples showed no recovery of the internal and surrogate standards: sample 7112-1523-N-L was a liquid, and the total volume collected was used in the initial analysis making reanalysis impossible; samples 7041-1192-N-L and 7037-1206-M-L were determined not to be analyzable after three separate attempts failed to achieve measurable recovery. Therefore, no results are reported for these three samples.

The initial result for sample 7129-1548-M-L was out of the dioxin calibration range and reanalysis of a one gram aliquot was performed. Other samples were initially analyzed as one gram aliquots; some reanalyses as extract dilutions were required. Table 5.3.6-2 summarizes the reanalysis results. These are reported in all cases because they are more representative of the sample aliquot analyzed.

5.4 SEWERS AND SUMPS

Sewer and sump samples were taken as described in the Subsection 4.2.4. Four sewer and eight sump samples were collected for dioxin analysis. Figure 4.2.4-1 indicates the sewer and sump locations on site. Table 5.4-1 summarizes the dioxin concentrations found in these samples and Table 5.4-2 presents the dioxin results for each specific location sampled.

Table 5.4-3 provides a summary of dioxin reanalysis results for the sewer and sump samples. Of the 12 samples taken, nine (75 percent) required reanalysis because the initial result was out of the dioxin calibration range. Because of the nonhomogeneity of the sludge samples collected and significant instrumental detector saturation, results for the one gram reanalysis and dilutions varied from the original analysis results. The reanalysis results are reported herein, because the analytical results are more representative of the aliquot analyzed. However,

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5.5.2 Analytical Laboratory Testing

Near-surface and boring soil samples were collected as described in Subsection 4.2.5. Each sample was uniquely identified in accordance with the protocol described in Subsection 4.1.2 and presented in Appendix A. The elevation codes used to designate samples at a specific location are schematically represented in Figure 5.5.2-1. Samples in the 100 series were taken in the fill; the 200 series designates the silt layer; and the 300 series samples were obtained from the glacial fluvial sand underlying the silt.

Near-surface soil samples were obtained to a depth of 60 inches. Samples from depth intervals of zero to 6 inches, 6 to 12 inches, and 12 to 24 inches were collected for the chemical analyses designated in the Work Plan. Below a depth of 24 inches, near-surface locations were continuously sampled at 12-inch intervals to a depth of 60 inches. Chemical analyses were not performed at samples from 24 to 60 inches as part of the current site investigation; however, the samples have been archived at the Director's Office Laboratory for possible future analysis. It is noted that prescribed building times preclude future analysis for all parameters except diatom. The archived near-surface soil samples are provided in Appendix E.

Boring soil samples were obtained from the silt layer. In accordance with the Work Plan, the samples were composited from Shelby tubes. To fully exploit the opportunity to collect additional near surface soil samples, the boring sampling program was expanded to include samples from zero to 24 inches in depth in increments of zero to 6 inches, 6 to 12 inches, and 12 to 24 inches.

Table 5.5.2-1 presents the analytical program conducted for the near-surface and boring soil samples. Chemical analysis was not performed for samples obtained below the silt layer.

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are not considered significant. In general, despite the stated difficulties associated with reanalysis, the results are considered more reliable than the initial analysis, and are reported for all samples where reanalysis was performed.

Table 5.5.2.1-3 summarizes the near-surface soil sample dioxin results by sample location and depth. Figures 5.5.2.1-1 through 5.5.2.1-3 present the dioxin results at depth intervals of zero to six inches, six to 12 inches, and 12 to 24 inches, respectively.

Priority pollutants were defined as the 157 compounds in the acid/base/neutral (semi-volatile), volatile organics, pesticides and PCB, herbicides, metals, total cyanide, and total phenol analyses plus the 40 extraneous peak searches. Forty-two near-surface soil samples were analyzed for priority pollutants. Table 5.5.2.1-4 presents the organic priority pollutant analysis levels and dilution factors applied to these samples. Reference is made to Appendix E for the definition of levels, and for the complete quantitative results.

Table 5.5.2.1-5 summarizes the detected base/neutral and acid organic compounds identified in the near-surface soil samples. Of the 69 semi-volatile compounds, 28 (41 percent) were identified one or more times in the depth intervals of zero to 6 and 12 to 24 inches. At zero to 6 inches, 24 compounds were identified. For 12 to 24 inches, 26 compounds were identified.

A summary of the detected volatile organic compounds in the near-surface soil samples is presented in Table 5.5.2.1-6. Of the 38 volatile organic compounds, 13 (34 percent) were identified one or more times. Methylene chloride and acetone were identified most frequently of the 13 compounds identified (42 of 42 and 28 of 42 samples, respectively). However, these concentrations are typically attributable to background levels due to handling either during collection, shipping, or in the

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5.5.2.2 Boring Soil Samples

Boring soil samples were collected at thirteen locations on site. Five samples were obtained for designated analyses at each of seven locations. At Boring B-2, a silt sample was not recovered, resulting in only four samples at that location. Table 5.5.2-1 indicates the designated chemical analyses for the boring soil samples and Table 5.5.2.2-1 summarizes the samples collected. Appendix E contains an inventory of the 124 samples archived for geotechnical purposes and possible future dioxin analysis. Also, Appendix E provides a summary of all quantitative analytical results and the results of the 40 extraneous peak searches for the boring soil samples.

Of the 39 boring soil samples analyzed for dioxin, 35 (90 percent) samples had identifiable dioxin concentrations. For nine of the 39 (23 percent) samples, reanalysis of smaller sample aliquots (1 gram) or dilutions were required to yield results within the instrumental calibration range. Table 5.5.2.2-2 presents a summary of the samples reanalyzed, the corrective actions taken, and the initial and reanalysis results. One one-gram sample required further dilution to produce results within the instrumental calibration range. Due to instrumental saturation and the inhomogeneity of soil samples, the reanalysis values increased as expected. In general, the values were of the same order of magnitude and reanalysis values were used for reporting purposes because they were more reliable than the initial analysis.

A summary of the dioxin results for the boring soil samples by sample location and depth is presented in Table 5.5.2.2-3. The results of dioxin analyses are also shown on the site cross sections in Figures 5.5.1-1 through 5.5.1-4. At depths of zero to 6 inches, the dioxin concentrations ranged from 19.7 ppb to 2700 ppb. At 6 to 12 inches, the dioxin concentrations ranged from 7.5 ppb to 3510 ppb, and at 12 to 24 inches, the dioxin concentration ranged from 4.7 ppb to 830 ppb. Samples from directly above the silt (Elevation Code 109) had dioxin

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at 12 to 24 inches, eight compounds were identified. For samples from above silt (Elevation Code 109), seven compounds were identified. Figure 5.5.2.2-3 presents the results for key semi-volatiles for samples from immediately above the silt layer.

A summary of the detected herbicides, pesticides, and PCB's in the boring soil samples is given in Table 5.5.2.2-7. Of the 35 herbicides, pesticides, and PCB compounds, 10 (29 percent) were identified one or more times in the samples from zero to 6 inches, 12 to 24 inches and above the silt. At zero to 6 inches, nine compounds were identified; at 12 to 24 inches, nine compounds were identified; and in the samples from above the silt, eight compounds were identified. Figure 5.5.2.2-4 presents the results for key pesticides and herbicides for samples from above the silt layer.

The detected inorganic parameters as defined by the 13 metals, total cyanides and total phenols in the boring soil samples are summarized in Table 5.5.2.2-8. Of the 13 metals, 11 (85 percent) were identified one or more times. At zero to 6 inches, ten metals were identified; at 12 to 24 inches, 11 metals were identified; and in the samples from above the silt, 11 metals were identified. Selenium and thallium were not identified. All samples had positive cyanide results and 7 out of 8 phenol results were positive.

The concentrations of selected priority pollutants are presented in Figures 5.5.2.1-4 through 5.5.2.1-7. Figures 5.5.2.2-3 and 5.5.2.2-4 present similar results for the boring soil samples from above the silt.

5.5.2.3 Additional Dioxin Analyses

Because dioxin was identified in the silt layer on site, an evaluation was initiated to investigate potential cross contamination, and/or instrumental carry over from previous injections. As part of the evaluation, the laboratory blanks and the chronological analysis order were

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As previously discussed the additional silt zone samples were not from the same depth as the initial samples. The evaluation of the data to define potential cross contamination during boring or sample collection was inconclusive.

5.6 GROUND WATER

The results of the ground water investigation at the site are presented in the following subsections. The results include monitoring well and piezometer levels, the determination of hydraulic conductivities from field slug test data, the calculation of ground water flow rates, and the chemical analysis data for the ground water samples.

5.6.1 Ground Water Levels

Ground water levels were observed on October 15, 1984 in the eight monitoring wells installed at the site (Figure 4.2.6-1). A summary of the observed levels is provided in Table 5.6.1-1. A comparison of monitoring well levels and Passaic River levels measured on this date is provided in Figures 5.6.1-1 through 5.6.1-8. As can be seen from the figures, the three monitoring wells nearest to the river (MW-1A, 2A, and 3A) fluctuate with the river. Ground water level contours based on these data are provided in Figure 5.6.1-9.

5.6.2 Hydraulic Conductivities

The slug test data were incrementally digitized from the continuous strip chart data taken in the field. This was done by selecting an arbitrary datum and scaling off values of time and pin deflection as a percent of full scale. Data reduction was accomplished using special forms to compute the value for head in feet for each selected time. The data were then input to the computer program SLUGT.

The program SLUGT computes aquifer transmissivity (T) and hydraulic conductivity (k) using two independent methods. The first method is the method of Cooper, Bredehoeft, and Papadopoulos (1967) and applies to

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As seen in Table 5.6.2-1, the representative mean hydraulic conductivity values for the most permeable zone range across the site from 3 to 200 feet per day. If the value of 200 feet per day at MW-2A is omitted, the range becomes 3 to 40 feet per day. Such ranges in mean permeability for the most permeable zone within a limited site area are to be expected when fill materials are involved. This is particularly true considering the nature of the fill--brick, building debris, wood, and other organic materials, in addition to the predominant sand, gravel, and varying amounts of silt.

5.6.3 Ground Water Flow Rates

Ground water flow rates were calculated based on the hydraulic conductivities presented in Table 5.6.2-1 and the gradients (change in head per unit distance) determined from Figure 5.6.1-9. An effective porosity of 0.30 was used for the fill. From the center of the site northward to the river, the computed ground water flow rate ranged from 0.6 to 4.0 feet per day. From the center of the site to the south, the range was 0.5 to 1.3 feet per day.

5.6.4 Analytical Laboratory Testing

Ground water samples were collected from each of the eight on-site monitoring wells as described in Subsection 4.2.6.3. Each sample was uniquely identified in accordance with the protocol described in Subsection 4.1.2 and presented in Appendix A. Sampling was conducted on October 9, 1984, and again on October 30, 1984, generating two sets of eight samples. Based on these preliminary dioxin results, the ground water from monitoring well MW-2A for a third time on December 14, 1984.

The first two rounds of ground water samples from all eight wells were analyzed for full priority pollutants plus 40 and dioxin. The third ground water sample from MW-2A was analyzed only for dioxin. Appendix F contains summaries of the quantitative analytical results (organic

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first round samples and six compounds (17 percent) were identified in the second round samples.

The detected inorganic parameters as defined by the 13 metals, total cyanide, and total phenols in the ground water are summarized in Table 5.6.4-6. Of the 13 metals, 11 (85 percent) were identified one or more times in the first round samples and 12 metals (92 percent) were identified in the second round samples. Seven of the eight ground water samples had positive cyanide results in each round. Total phenols analysis showed positive results for all 16 samples analyzed.

5.7 PASSAIC RIVER WATER

5.7.1 Passaic River Levels

The mean tidal range (difference in height between mean high water and mean low water) is reported by the National Oceanic and Atmospheric Administration as 5.1 feet. The spring range (average semi-diurnal range occurring semi-monthly as a result of the full moon and new moon) is reported to be 6.1 feet. Tide levels in the Passaic River were measured over one complete tidal cycle on October 15, 1984, and were compared with ground water levels in the on-site monitoring wells. This information is presented in Figures 5.6.1-1 through 5.6.1-8.

5.7.2 Analytical Laboratory Testing

Two Passaic River water samples were collected (one on October 9, 1984 and one on October 30, 1984), concurrent with the ground water sampling for the eight on-site wells. These samples were collected by the procedures described in Subsection 4.2.7.2. Each sample was uniquely identified in accordance with the protocol described in Subsection 4.1.2 and presented in Appendix A.

Both samples were analyzed for dioxin and full priority pollutants. Priority pollutants were defined as the 157 compounds in the acid/base/

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the dioxin concentrations ranged from 0.53 to 10.8 ppb with six samples having nondetectable dioxin concentrations at a detection limit of 0.78 ppb. At 12 to 24 inches, the dioxin concentrations ranged from 0.63 to 130 ppb with four samples having nondetectable dioxin concentrations at the 0.78 ppb detection limit.

Additional river sediment samples were obtained to a depth of six feet at Station 1-3-0. A geotechnical log is provided in Appendix B, and shows the sample locations. Table 5.8-3 presents the dioxin results of samples obtained at Station 1-3-0.

Priority pollutants were defined as 157 compounds in the acid/base/neutral (semi-volatile), volatile organic compounds (VOC), pesticides and PCB's, herbicides, metals, total cyanides, and phenols analyses plus 40 extraneous peak searches. The organic priority pollutant analysis levels and dilution factors that were utilized in the analyses of the sediment samples are presented in Table 5.8-4. Complete quantitative results for each sample are given in Appendix G.

A summary of all the detected base/neutral/acid organic compounds identified in the Passaic River sediments is given in Table 5.8-5. Of the 69 semi-volatile compounds, 17 (25 percent) were identified one or more times in the zero- to 12-inch or 12- to 24-inch samples. Fourteen compounds were identified in 28 percent of the samples at zero to 12 inches. Seventeen compounds were identified in 36 percent of the samples at 12 to 24 inches.

A summary of all the detected volatile organic compounds identified in the Passaic River sediments is presented in Table 5.8-6. Of the 38 volatile organic compounds, 10 were identified in one or more samples at the zero- to 12-inch or 12- to 24-inch depths. Eight compounds were identified in 33 percent of the samples at depths of zero to 6 inches. Ten compounds were identified in 46 percent of the samples at depths of 12 to 24 inches.

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Appendix H contains the quantitative results of the priority pollutant analysis. Of the 69 acid/base/neutral compounds, 20 (24 percent) were reported one or more times in the three samples analyzed. For the three samples analyzed, the results were as follows: of the 38 volatile organic compounds, three (8 percent) were reported one or more times; of the 35 herbicide, pesticide, and PCB compounds, two were reported one or more times; and of the 13 metals, 11 were reported all three times.

5.9.2 Newark

Samples collected at Harrison Avenue, Raymond Boulevard, and Roanoke Avenue were taken to establish a background for the area. Figure 4.2.9-1 indicates the sampling locations. Three samples were collected at depths of zero to 6 inches and analyzed for priority pollutants and dioxin. All quantitative results for each sample are given in Appendix H. The dioxin results of the three background samples from Newark are presented in Table 5.9.2-1.

A summary of the base/neutral/acid compounds detected in the near-surface soil samples from the site and the Newark background locations is presented in Table 5.9.2-2. Of the 69 semi-volatile compounds, 16 were identified in the Newark background samples one or more times. A summary of the volatile organic compounds detected in the near-surface soil samples from the site and Newark background samples is presented in Table 5.9.2-3. Methylene chloride was the only compound reported. A summary of the herbicides, pesticides, and PCB compounds detected in the near-surface soil samples from the site and the Newark background samples is provided in Table 5.9.2-4. Of the 35 herbicide, pesticide, and PCB compounds, three were detected one or more times. A summary of the inorganic parameters detected in the near-surface soils for the site and the Newark background samples is presented in Table 5.9.2-5. Of the 11 metals, 11 were identified one or more times. Positive total cyanide and phenol results were reported for four of the six samples analyzed.

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Other designations are used for specific parameters. These are:

- o Solubility - The ability of the material to combine with water, and reported on a visually determined scale of none, slight, moderate, and complete. The designation is subjective.
- o Temperature Change - As the sample material was mixed with water, the relative temperature of the reaction vessel was assessed by touch and reported as increasing, decreasing, or no change (none detected).
- o Percent Lower Explosive Limit (LEL) - With the detection meter calibrated to 10 percent, vapor from the sample was tested for percent of combustible organic vapors present. Only four composite samples (Pit-2, PP-2, PP-3, PP-6) and one individual drum (No. 120) registered on the LEL meter. Sample PP-3 registered 6 percent, PP2 and PP-6 registered 3 percent, the two Pit-2 drums showed 2 percent, and Drum 120 registered 2 percent.
- o Oxidizable Material (OX) - This test was performed to determine the presence of any material that readily initiates the combustion of organic matter. All of the individual drums or composite samples showed negative tests for the presence of oxidizable materials.
- o Peroxides - All tests for the presence of peroxides produced negative results.
- o Sample Type - This visual classification of the physical state of the sample was presented simply as solid, liquid, sludge, or multilayered.
- o Open-Cup Ignitability - This test identifies wastes which present fire hazards from ignitability under conditions existing during routine storage, disposal, and transportation. Results were shown as: does not ignite when heated (DNI) or ignites when heated (IWH). Other assessments were also made as to the ability to sustain combustion and the type of flame generated.
- o Open Cup Flashpoint - The material was heated in an open cup to 140 degrees Fahrenheit. If it ignited the temperature was recorded in degrees

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analyzed for 2,3,7,8-tetrachlorodibenzofuran (2,3,7,8-TCDF) and octa-chlorodibenzo-p-dioxin (OCDD). The 2,3,7,8-TCDF analyses were run under the same conditions as the dioxin (2,3,7,8-TCDD) analyses. This includes the seven isomer dioxin performance mixture for demonstrating separation of the dioxin isomers. An analogous mixture for all of the furan isomers is not available, because TCDF reference standards are not available. It has been reported in the scientific literature that the TCDF isomers will be separated under the same conditions that separate the TCDD isomers; therefore, the 7 isomer TCDD mix was used to demonstrate this capability. However, since the full separation of the TCDF isomers cannot actually be demonstrated, the result reported for 2,3,7,8-TCDF should be considered the maximum concentration for this compound in the sample. Procedures for analysis of 2,3,7,8-TCDF and OCDD were presented in the Work Plan.

A total of 156 soil samples were analyzed for dioxin, including additional analyses and background samples. Twenty-four samples were analyzed for 2,3,7,8-TCDF and OCDD. Table 5.11-1 presents the results of these analyses.

5.12 ANALYTICAL RESULTS FOR QUALITY ASSURANCE/QUALITY CONTROL CHECKS

Quality assurance/quality control checks were performed routinely throughout the course of the sampling and analysis activities. Several levels of QC checks were implemented, as described in the QA Plan contained in the Work Plan, including field/trip blanks, individual laboratory analysis-specific QC measures, an additional set of "Program QC" samples initiated by the QA Program Director, and specific QC samples initiated by the NJDEP.

Precision, accuracy, and completeness objectives for the analytical program were established in the Work Plan, and are presented in Table 5.12-1. Similar QC acceptance criteria were available for those analyses performed under EPA's CLP protocol. Where available, these analyte specific acceptance limits were used.

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2.7 ng. All samples collected on September 7, 1984 prior to preparation of the wipe blank showed no detectable dioxin; therefore, the one wipe sample associated with it was scheduled for recollection (at an adjacent location), with a new field blank. The new field blank showed no detectable dioxin. On September 20, all four collected wipe samples indicated extremely high levels of dioxin; therefore the blank result was negligible.

The two chip field blanks and the paired trip blanks showed no detectable levels of dioxin.

5.12.2 Individual Laboratory Quality Control Checks

Following are discussions of the QC checks performed by the three ITC laboratories which provided analyses.

5.12.2.1 Organic Priority Pollutant Analyses - ITC Cerritos Laboratory

For all parameters (except herbicides) EPA-Contract Laboratory Program (CLP) procedures were followed, as described in the Work Plan. EPA Method SWA 8150 was used for the chlorinated herbicide analyses; these compounds are not included in the CLP protocols, and are, therefore, not subject to the same QC criteria.

Instrument calibrations were performed and documented every eight hours. Support data for these QC checks are contained in the complete laboratory batch reports provided with each set of sample results. Laboratory blanks were also analyzed at least once every eight hours for each type of analysis performed. No significant contamination was observed in the laboratory blanks for extractable parameters (BNA, pesticide, herbicide). Volatile blanks did consistently show low levels of methylene chloride--generally below the quantitation limit. Complete data for all laboratory blanks are also contained in the batch reports.

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for each surrogate is summarized in Table 5.12.2.1.1-2. All of the pesticide surrogate outside recoveries, in low and medium level analyses, are due to dilution of the extracts prior to analysis. Recoveries were not obtained for these samples.

One soil sample showed very low recoveries for all six BNA surrogates; three of these were outside the QC limits. In accordance with CLP protocol, the sample was reextracted; reanalysis showed identical surrogate recovery results, indicating a significant matrix effect for this sample.

As presented in Table 5.12.2.1.1-1, all average percent recoveries for surrogate compounds are well within the stated QC acceptance criteria (as defined in Table 5.12.2.1-1) for soil samples. The consistency of the recovery results, as measured by the standard deviation, is also acceptable for each compound.

Matrix Spike/Matrix Spike Duplicates

A total of 75 soil matrix samples were analyzed for all organic priority pollutants. Table 5.12.2.1.1-3 summarizes the number of MS/MSD pairs analyzed by analysis fraction and level. All QC check frequencies for soils are greater than the minimum 5 percent objective.

Tables 5.12.2.1.1-4 through 5.12.2.1.1-6 summarize the average percent recoveries and RPD's for all spike compounds. Recoveries outside QC acceptance limits are not included in these calculations. Excluding pesticide spikes, only 4 percent of the results were outside of recovery limits. These all occurred in the medium BNA analysis of a single MS/MSD pair. The compounds were pyrene and phenol. Because the recoveries properly duplicated for both compounds in the MS and MSD, the RPD fell in the acceptable range.

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Matrix Spike/Matrix Spike Duplicates

A total of 15 sediment samples were analyzed for full organic priority pollutants; one MS/MSD pair was analyzed for each analysis fraction and level, as outlined below:

Low VOA	- 1 of 15 samples	6.7% frequency
Low BNA	- 1 of 10 samples	10% frequency
Med BNA	- 1 of 5 samples	20% frequency
Low Pesticide	- 1 of 13 samples	7.7% frequency
Med Pesticide	- 1 of 2 samples	50% frequency
Herbicide	- 1 of 15 samples	6.7% frequency

The 5 percent minimum QC check frequency was achieved.

Tables 5.12.2.1.2-2 through 5.12.2.1.2-4 summarize the average percent recoveries and RPD's for all spike compounds. Recoveries outside QC acceptance limits are not included in these calculations; however, only two were noted, and these were for pyrene in the MS and MSD pair analyzed. The observed recoveries for pyrene were 20 percent in each QC sample, only slightly below the lower QC acceptance limit of 26 percent.

The accuracy (average percent recovery) and precision (RPD) measures for all priority pollutant spike compounds are well within the stated QC acceptance limits. Herbicide spike recoveries range from 47 to 95 percent recovery except for Dinoseb, which shows an average recovery of 2.1 percent.

Sediment priority pollutant data quality has not been affected by the small number of results outside of QC objectives.

5.12.2.1.3 Water QC Summary

Surrogate Recoveries

Table 5.12.2.1.3-1 summarizes the average surrogate recovery results for all water samples (all were performed as low-level analyses). Recoveries outside of QC limits are not included. The number of results

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Middlebrook Pike laboratory. EPA methods, as specified in the Work Plan, were used for analysis.

Calibration standards and laboratory blanks were analyzed and documented a minimum of once per eight-hour shift. Contamination of sufficient concentration to affect results was not observed in any of the laboratory-prepared blanks. Full documentation of all standards, blanks, and daily control charts is contained in the laboratory files.

Table 5.12.2.2-1 summarizes the spike and duplicate analyses performed for each sample matrix. In all cases, QC checks were greater than 5 percent of the total samples analyzed for each matrix.

The results of spike recovery, laboratory blanks, and duplicate (blind split) analyses are contained in Appendix J. Table 5.12.2.2-2 summarizes the RPD values for inorganic duplicates on water and soil/sediment matrices. Nine of the RPD values are greater than 20 percent, indicating relatively low reproducibility between original and duplicate sample results. However, as Figure 5.12.2.2-1 shows, the method itself is precise to within 10 percent for standard response factors over a month-long period. Two explanations could account for the observed RPDs: (1) many of the measurements are at or near the detection limit, where the method is, by definition, at the limit of its precision and accuracy; and (2) sample nonhomogeneity and/or matrix effects make it difficult to reproduce results precisely, regardless of the degree of homogenization.

The average percent recovery for each spike compound, including all matrices, is shown in Table 5.12.2.2-2.

All of these average recoveries are within the QA objective limits as stated in the Work Plan.

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Table 5.12.2.3-2 presents the spike recovery data for the QC samples. Thirteen of the 27 samples selected for QC analysis showed dioxin at concentrations greater than or equal to 10 ppb; the spike level of 1.0 ppb becomes negligible for these samples, negating meaningful recovery calculations. These values have, therefore, been excluded from this table. The average percent recovery is calculated to be 93 (± 17) percent.

Partial scan GC/MS confirmations were performed on 35 samples with positive dioxin results. Of these, 29 (83 percent) passed all criteria required for a final confirmation of the presence of 2,3,7,8-TCDD. Two of the four failures were due to dioxin levels too low to be detected by the partial scan method. Back-up documentation of these analyses is contained in the dioxin laboratory batch reports.

Two special QC samples were analyzed in an effort to determine the efficiency of the wipe sampling technique used. A piece of aluminum foil, measuring 46 x 54 cm (2,480 cm²), was spiked with 250 μ l of a 0.2 ppm 2,3,7,8-TCDD standard solution (equivalent to 50 ng dioxin) by applying the solution in an "S" pattern across the foil surface. The spiked foil was allowed to air dry for 30 minutes, until no solvent was observed. A 3-by-3-inch gauze wipe was wetted with 10 ml of hexane, and then used to "sample" the foil surface in accordance with the wipe sampling procedure used on site. This spiked foil sample was prepared, sampled, and extracted in duplicate at the Directors Drive laboratory.

Results for the duplicate samples were as follows:

SAMPLE NO.	AMOUNT OF DIOXIN RECOVERED	% RECOVERY
J2196	39.8 ng	80
J2197	35.4 ng	71

The average recovery is 76 (± 4.5), which is within the acceptable range of precision for the dioxin analysis method.

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No significant contamination was detected in the wipe blanks or the water blank as shown in Table 5.12.3.1.-1 and Appendix J. The soil blanks were aliquots of a supply of Clarksburg soil, originally obtained from the EPA-OHMSETT facility in Leonardo, New Jersey and described as "virgin with respect to 2,3,7,8-TCDD." The soil had been stored, however, in the dioxin preparatory laboratory since its arrival in 1983; thus, contamination from its storage location most likely explains the low levels of dioxin detected. The average of 1.1 (± 0.4) ppb is just above the detection limit for the analysis. The water blank result was below the instrumental detection limit.

5.12.3.2 QC Blank Spikes

Table 5.12.3.2-1 provides the recovery results for the QC Program spiked wipes and soils. The average percent recovery for the wipe samples is 90 (± 4.2). For the soils, the average recovery is 118 (± 9.5). Both of these averages are within acceptable limits, as prescribed in Table 5.12-1.

5.12.3.3 Directed Sample Splits

Table 5.12.3.3-1 contains the duplicate results summary for dioxin in the 10 sample splits which were initiated at the QA Program Manager's direction. Complete parameter results for the three soil and one water split designated for full analysis are presented in Appendix J.

For the dioxin results in soil samples, a major difference is noted in the RPD values for comparison of 10-gram aliquot samples versus one-gram aliquot samples. In all four cases, the RPD is vastly improved for the 10-gram results. This is most probably due to the difficulty encountered in obtaining a one-gram aliquot that is truly representative of the entire sample, regardless of the degree of homogenization. The one-gram RPD values have, therefore, been excluded from calculation of the average, which is 33 (± 34). This is within acceptable QC limits for overall precision.

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5.12.4 NJDEP-Designated Quality Control Checks

The NJDEP On-Scene Coordinator(s) (OSC) initiated two types of quality control samples: (1) collection of sample splits in the field, for analysis at an independent laboratory in addition to the ITC laboratories, and (2) assignment of soil proficiency samples for dioxin analysis.

Table 5.12.4-1 lists the samples split at the time of collection at the direction of the OSC. ITC results for these samples, for the analyses indicated, are reported herein with all other analytical results. Results for samples analyzed at NJDEP's chosen independent laboratory have not been received by ITC, and are, therefore, not included in this report.

Table 5.12.4-2 contains the dioxin results for the NJDEP soil proficiency samples. One sample (not listed in the table), ITC No. 9400-1763-T-L (NJDEP A037), was received broken at the laboratory on October 23, 1984; the sample was voided as instructed by L. Geiger (NJDEP-OSC); therefore, analysis was not required. All samples labeled "blank spike" were described as "blank-to be spiked with 2,3,7,8-TCDD at 1 ppb" on the Chain-of-Custody record. The confirmed concentration of the spiking solution used was 160 pg/ul; 100 ul is spiked onto 10 grams of soil prior to extraction. Thus, the actual spike level used for these samples was 1.6 ng/g, or 1.6 ppb.

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5.0 REFERENCES

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TABLES

TABLE 5.1-1
AMBIENT AIR SAMPLES
RESULTS: IRON AND MANGANESE
(10-31-84)

COLLECTION DATE		RESULTS (ug/filter)			NJDEP SAMPLE DATES	SAMPLES TO BE ANALYZED FOR FULL AMBIENT AIR PARAMETERS
		Fe	Mn	Fe+Mn		
09-08-84	A002-0142-A-K	252	7.1	259		
09-09-84	A002-0143-A-K	704	21	725	X	
09-10-84	A002-0144-A-K	1,280	34	1,314*		0
09-11-84	A002-0181-A-K	1,550	41	1,591*		0
09-12-84	A002-0182-A-K	1,740	91	1,831*		0
09-13-84	A002-0183-A-K	917	31	948		
09-14-84	A002-0410-A-K	823	22	845		
09-15-84	A002-0412-A-K	259	7.1	266	X	
09-16-84	A002-0413-A-K	702	17	719		
09-17-84	A002-0414-A-K	1,230	34	1,264*		0
09-18-84	A002-0415-A-K	1,090	31	1,121*		
09-19-84	A002-0597-A-K	1,770	57	1,827*		0
09-20-84	A002-0598-A-K	992	34	1,026		
09-21-84	A002-0711-A-K	1,080	38	1,118	X	0
09-22-84	A002-0712-A-K	676	21	697		
09-23-84	A002-0713-A-K	702	22	724		
09-24-84	A002-0714-A-K	1,520	37	1,557*		0
09-25-84	A002-0843-A-K	2,030	63	2,093*		0
09-26-84	A002-0844-A-K	745	19	764		
09-27-84	A002-0845-A-K	810	19	829	X	
09-28-84	A002-1072-A-K	566	15	581		
09-29-84	A002-1073-A-K	448	11	459		
09-30-84	A002-1074-A-K	597	25	622		
10-01-84	A002-1082-A-K	312	8.5	320		
10-02-84	A002-1083-A-K	590	26	616		
10-03-84	A002-1084-A-K	1,200	34	1,234*	X	0
10-04-84	A002-1241-A-K	1,460	52	1,512*		0
10-05-84	A002-1242-A-K	1,030	21	1,051		
10-06-84	A002-1243-A-K	1,040	27	1,067		
10-07-84	A002-1329-A-K	610	19	629		
10-08-84	A002-1330-A-K	1,160	37	1,197*		

* Denotes 10 highest total (Fe + Mn) results.

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DATE AND SAMPLE ID	PARAMETERS	Particulates: (ug/m ³)					Metals: (ug/m ³)							
		TSP	IPM				Iron	Manganese	Cadmium	Copper	Nickel	Lead	Vanadium	Zinc
9-10-84	-0144-A-K	108	118	79	0.908	0.732	0.019	0.024	0.003	0.031	0.019	0.364	0.023	0.240
9-11-84	-0181-	117	74	0.997	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.399	0.017	0.321
9-12-84	-0182-	102	165	85	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.423	0.041	0.368
9-17-84	-0414-	102	165	85	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.298	<0.014	0.227
9-19-84	-0597-	100	54	123	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227
9-21-84	-0711-	85	153	123	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227
9-24-84	-0714-	153	254	196	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227
9-25-84	-0843-	254	196	56	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227
10-3-84	-1084-	114	56	87	0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227
10-4-84	-1241-	87	75		0.944	1.004	0.026	0.052	0.003	0.033	0.025	0.348	0.016	0.227

TABLE 5.1-2
SITE INVESTIGATION
AMBIENT AIR RESULTS FOR
TOTAL SUSPENDED PARTICULATES (TSP)
INHALEABLE PARTICULATE MATTER (IPM) AND METALS

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TABLE 5.1-3
SITE INVESTIGATION
AMBIENT AIR RESULTS FOR DIOXIN

DATE AND SAMPLE ID PARAMETERS	9-10-84 94-0144-A-X	9-11-84 94-0181-	9-12-84 94-0182-	9-17-84 94-0418-	9-19-84 94-0597-	9-21-84 94-0711-	9-24-84 94-0714-	9-25-84 94-0863-	10-3-84 94-1084-	10-4-84 94-1261-
Dioxin (pg/m ³)	86	MD(<8)	MD(<6)	MD(<31)	MD(<8)	MD(<4)	286	MD(<10)	MD(<6)	MD(<15)

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TABLE 3.1-4

DATE AND SAMPLE ID	PARAMETERS	Vinyl Chloride	VOC's: (ug/m ³)	Methylene chloride	1,1-dichlorobenzene	Chloroform	1,2-dichloroethane	1,4-dioxane	Carbon tetrachloride	Trichloroethene	Benzene	1,1,2-trichloroethane	1,2-dibromethane	Tetrachloroethene	1,1,2,2-tetrachloroethane	Toluene	Chlorobenzene	Ethylbenzene	Styrene	m-Xylene	p-Xylene	o-Xylene	o-chlorotoluene	p-chlorotoluene	o-dichlorobenzene	p-dichlorobenzene
9-10-84	-0144-A-K	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-11-84	-0181-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-12-84	-0182-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-17-84	-0414-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-19-84	-0597-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-21-84	-0711-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-24-84	-0714-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
9-25-84	-0843-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
10-3-84	-1084-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08
10-4-84	-1241-	MD	0.15	0.26	0.33	<0.12	MD	2.93	2.74	0.94	2.27	MD	0.53	0.55	MD	0.28	0.30	3.10	MD	0.56	1.08	1.03	1.08	1.08	1.08	1.08

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DATE AND SAMPLE ID	PARAMETERS	Asbestos (fibre) cm ³
9-10-84	-0144-A-E	0.009
9-11-84	-0181-	0.002
9-12-84	-0182-	0.001
9-17-84	-0414-	0.004
9-19-84	-0597-	0.003
9-21-84	-0711-	0.002
9-24-84	-0714-	0.003
9-25-84	-0843-	0.005
10-3-84	-1084-	0.001
10-4-84	-1241-	0.003

TABLE 5.1-5
SITE INVESTIGATION
AMBIENT AIR RESULTS FOR ASBESTOS

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* - higher detection limit due to sample matrix interference.

DATE AND SAMPLE ID	PARAMETERS	Pesticides (ng/m ³)	Benzene sulfon- chloride	Tetrachlorobenzene	4-chlorobenzene sulfon chloride	4-methoxybenzene sulfon chloride	Hexachlorobenzene	2,4,5-Trichloro ester)	Oven	p,p'-DDE
9-10-84	-0144-A-K	2.97	<2.38	<2.22	<14.39*	<2.55	<14.39*	<7.55*	<2.36	<2.44
9-11-84	-0181-	<0.10	<0.10	<12.7	<14.39	<0.06	<0.75*	1.74*	<0.10	<1.06*
9-12-84	-0182-	<13.5	<13.6	<12.7	<14.39	<0.06	<0.75*	1.74*	<0.10	<1.06*
9-17-84	-0414-	43.92	29.93	50.79	<71.9*	47.77	39.57	49.06	50.68	18.52
9-24-84	-0714-	0.71	0.32	0.79	0.97	0.32	0.47	0.72	<0.07	0.30
9-25-84	-0843-	3.38	2.52	2.70	2.19	2.36	1.80	2.60	2.94	1.18
10-3-84	-1084-	<1.35	<1.36	<1.27	<1.44	<1.51	<1.35	<1.35	<1.35	<1.39
10-4-84	-1241-	4.73	4.08	<3.17	<3.60	<3.18	<3.60	3.77	<3.38	<3.37

TABLE 5.1-6
SITE INVESTIGATION
AMBIENT AIR RESULTS FOR PESTICIDES

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TABLE 5.1-7
SITE INVESTIGATION
AMBIENT AIR RESULTS FOR PM10

DATE AND SAMPLE ID	PARAMETERS	PM ₁₀ (ng/m ³)							
9-10-84	-0144-A-K	0.51	0.63	1.01	0.80	1.41	0.72	1.24	1.35
		Benz(e)(h)flworanthene							
9-11-84	-0181-	0.54	0.67	3.01	0.90	2.50	1.66	2.28	1.10
		Benzo(a)pyrene							
9-10-84	-0144-A-K	0.70	1.08	2.10	1.96	2.35	2.10	1.09	2.55
		Benzo(g,h,i)perylene							
9-10-84	-0144-A-K	0.42	0.33	0.71	1.08	1.97	1.07	0.94	1.56
		Indeno(1,2,3-c,d)pyrene (Coronene)							
9-11-84	-0181-	0.68	0.38	0.09	0.28	1.07	0.46	0.89	0.72
		Phenanthrene							
9-10-84	-0182-	0.38	0.36	1.50	0.38	0.86	0.31	0.68	0.80
		Triphenylene							
9-11-84	-0182-	1.80	0.86	1.62	0.97	1.97	0.91	1.66	1.55
		Benzo(b)flworanthene							
9-10-84	-0182-	0.06	0.05	0.21	0.03	0.17	0.06	0.11	0.09
		Anthracene							
9-10-84	-0182-	1.36	1.18	6.66	1.01	2.69	1.07	2.26	2.01
		Flworanthene							
9-11-84	-0181-	1.19	1.16	1.76	1.13	2.42	1.47	1.86	2.01
		Pyrene							
9-10-84	-0144-A-K	0.78	0.65	0.56	0.76	1.45	0.59	1.22	1.28
		Benzo(a)anthracene							
9-10-84	-0144-A-K	2.05	2.34	2.90	2.79	*	1.27	2.72	2.59
		Benzo(a,h)anthracene							
9-10-84	-0144-A-K	0.73	0.63	0.54	0.58	1.39	0.53	1.12	1.32
		Chrysene							
9-10-84	-0144-A-K	7.16	5.75	*	6.05	12.02	5.70	11.99	10.40
		Perylene							

TABLE 5.2.1-1
INDUSTRIAL HYGIENE 2,3,7,8-TCDD MONITORING RESULTS

SAMPLE DESCRIPTION	RESULTS
IH-glass fiber filter: personnel sample	ND (0.41 ng/m ³)
IH-glass fiber filter: hi vol, clean area sample	ND (0.20 ng/m ³)
IH XAD2: backup to L0104	ND (0.27 ng/m ³)
IH-glass fiber filter: blank	ND (0.80 ng/sample)
IH-XAD-2 tube field blank	ND (0.75 ng/sample)
IH-glass fiber filter-personnel sample	ND (1.6 ng/m ³)
IH-GFF/XAD: 37-Cl sample spike	ND (1.2 ng/m ³)
IH-glass fiber filter-Hi vol, btwn tanks, process building	ND (1.0 ng/m ³)
IH-XAD: backup to L0161	ND (0.08 ng/m ³)
IH-glass fiber filter-blank	ND (1.1 ng/sample)
IH-XAD-blank	ND (0.36 ng/sample)
IH-glass fiber filter-personnel sample	ND (0.34 ng/m ³)
IH-glass fiber filter-field blank	ND (1.1 ng/sample)
IH-glass fiber filter-personnel sample	ND (0.10 ng/m ³)
IH-personnel sample: drum sampler assistant	ND (0.55 ng/m ³)
IH-personnel sample: driller (glass fiber filter)	ND (0.14 ng/m ³)
IH-glass fiber filter: btwn tanks and process building	ND (0.16 ng/m ³)
IH-XAD: backup to L0276	ND (0.37 ng/m ³)
IH-glass fiber filter-field blank	ND (0.11 ng/sample)
IH-XAD: field blank	ND (1.1 ng/sample)
Glass fiber filter: personnel	ND (0.12 ng/m ³)
Glass fiber filter: personnel	ND (0.91 ng/m ³)
Glass fiber filter: personnel	ND (0.99 ng/m ³)
Glass fiber filter field blank	ND (0.41 ng/sample)
IH-glass fiber filter-area decon	ND (0.16 ng/m ³)
IH-XAD: backup to L0444	ND (0.55 ng/m ³)
IH-glass fiber filter-personnel	0.74 ng/m ³
IH-glass fiber filter-field blank	ND (0.05 ng/sample)
IH-XAD2-field blank	ND (0.32 ng/sample)
IH-glass fiber filter-personnel	ND (0.31 ng/m ³)
IH-glass fiber filter-field blank	ND (1.0 ng/sample)
IH-glass fiberfilter-personnel, chip sampler	1.2 ng/m ³
IH-glass fiber filter-personnel, driller	ND (0.26 ng/m ³)
IH-glass fiber filter-field blank	ND (0.61 ng/sample)
IH-GFF/XAD: 37-Cl blank spike	ND (1.3 ng/sample)
IH-glass fiber filter-personnel, driller	ND (0.49 ng/m ³)
IH-glass fiber filter-personnel, tank sampling	ND (0.74 ng/m ³)
IH-XAD: backup to L0807	ND (0.55 ng/m ³)
IH-glass fiber filter-field blank	ND (0.51 ng/sample)
IH-XAD: field blank	ND (0.68 ng/sample)

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TABLE 5.2.1-1
(Continued)

SAMPLE DESCRIPTION	RESULTS
IN-glass fiber filter area	ND (1.01 ng/m ³)
IN-XAD2, area sample	ND (0.27 ng/m ³)
IN-glass fiber filter field blank	ND (0.5 ng/sample)
IN-XAD2 field blank	ND (0.55 ng/sample)
IN-glass fiber filter, personnel-drum crew	ND (2.02 ng/m ³)
IN-glass fiber filter, personnel-tank crew	ND (0.33 ng/m ³)
IN-glass fiberfilter, personnel-drillers	ND (2.9 ng/m ³)
IN-glass fiber filter-field blank	ND (0.4 ng/sample)
IN-glass fiber filter area	ND (0.28 ng/m ³)
IN-XAD2, area sample	ND (0.97 ng/m ³)
IN-XAD2, field blank	ND (2.7 ng/sample)
IN-glass fiberfilter, personnel-drums	ND (0.39 ng/m ³)
IN-glass fiber filter, personnel-tanks	ND (2.4 ng/m ³)
IN-glass fiber filter, field blank	ND (0.85 ng/sample)
IN-wipe: dcn line where personnel untape	52.2 ng/wipe
IN-wipe: stblz cloth in decon btwn brk area and D trlr	ND (16.4 ng/m ²)
IN-vater: final rinse tub in decon line	0.02 ppb
IN-wipe: stblz cloth in frnt of sampl trlr steps	ND (26.8 ng/m ²)
IN-Glass fiber filter, personnel-tank	ND (0.49 ng/m ³)
IN-glass fiber filter, field blank	ND (0.93 ng/sample)
IN-wipe: frm sampl head of instrmt No. 15084 after decon	ND (11.2 ng/m ²)
IN-wipe: from ostd body of decon instrmt No. 15084	ND (4.8 ng/m ²)
IN-wipe: field blank	ND (3.8 ng/wipe)
IN-glass fiber filter, personnel, tank	26.3 ng/m ³
IN-glass fiber filter, personnel, tank	ND (4.5 ng/m ³)
IN-glass fiber filter, field blank	ND (11.1 ng/sample)
IN-wipe: wheel of drill rig after decon	ND (4.4 ng/m ²)
IN-wipe: back of drill rig on steel plates after decon	84. ng/m ²
IN-wipe, field blank	ND (1.5 ng/wipe)
IN-glass fiber filter-frnt personnel drl rig dcn	98.9 ng/m ³
IN-glass fiber filter-bck personnel drl rig dcn	8.9 ng/m ³
IN-glass fiber filter-field blank	ND (3.2 ng/sample)
IN-glass fiber filter-area onstd dcn tent-drl rig	ND (18.1 ng/m ³)
IN-XAD2-area ostd decon tent for drill rig	ND (9.6 ng/m ³)
IN-XAD2- field blank	ND (52 ng/sample)
IN-glass fiber filter, personnel-soil crew	1.8 ng/m ³
IN-glass fiber filter, personnel-soil crew	1.7 ng/m ³
IN-wipe: bck of drl rig-dck stl plr-right side	72 ng/m ²
IN-wipe: steel high pressure air bottle	18 ng/m ²
IN-wipe: MSA air hose	124 ng/m ²
IN-wipe: steam jenny heater tower	8.4 ng/m ²
IN-wipe-field blank	ND (4.1 ng/wipe)

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TABLE 5.2.3-1
SUMMARY OF ORGANIC COMPOUNDS, ASBESTOS, AND
SULFURIC ACID RESULTS IN
INDUSTRIAL HYGIENE MONITORING SAMPLES

PARAMETER	CONCENTRATION RANGE	NUMBER OF POSITIVE RESULTS	NUMBER OF SAMPLES ANALYZED
2,4-D	ND	0	1
2,4,5-T	ND	0	1
Asbestos	0-0.09 fibers/cm ³	1	2
Sulfuric Acid	0.022 mg/m ³	1	1
Toluene ⁽¹⁾	0.097-0.099 mg/m ³	3	4
1,2-Dichloroethane	0.099 mg/m ³	1	4
Tetrachloroethane	1.44 mg/m ³	1	4
1,1,1-Trichloroethane	1.98 mg/m ³	1	4
Trichloroethylene	1.08 mg/m ³	1	4
1,1,2-trichloro-1,2,2-trifluoroethane	8.1 mg/m ³	1	4

(1) Toluene was also detected in field blank sampling tubes at comparable levels.

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TABLE 5.3-1
SUMMARY OF ASBESTOS ANALYSIS
BUILDINGS AND STRUCTURES

LOCATION	NUMBER OF SAMPLES	NUMBER OF POSITIVE RESULTS	RANGE OF RESULTS IN (%) BY ASBESTOS FORM	
			CHRYSTOTILE	AMOSITE
Lab	4	3	<1-26	-
Warehouse	3	1	25	4
Manufacturing Building	2	1	15	-
Process Building	3 ⁽¹⁾	2	14	5
Tank Farm	2	2	16	6.5
TOTALS	14	9	<1-25	4-6.5

(1) One sample lost in transit.

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TABLE 5.3-2
ASBESTOS ANALYSIS RESULTS
BUILDINGS AND STRUCTURES

SAMPLE NUMBER	DESCRIPTION	RESULTS (X)
<u>MANUFACTURING BUILDING</u>		
3100-1306-B-C	Mftg. Bldg., First Floor, Piping Insulation	Chrysotile 15
3100-1307-B-C	Mftg. Bldg., Boiler Room, Piping Insulation	None
<u>PROCESS BUILDING</u>		
4100-1305-B-C	Process Building, First Floor, Piping Insulation	Chrysotile 14
4100-0593-B-C	Process Building, First Floor, Piping Insulation	Amosite 5
4100-0594-B-C	Process Building, First Floor, Vessel Piping Insulation	Sample Lost
<u>TANK FARM</u>		
9500-1304-B-C	Northwest Tank Farm Piping Insulation	Amosite 6.5
9500-1308-B-C	Southeast Tank Farm Piping Insulation	Chrysotile 16
<u>WAREHOUSE</u>		
2100-0281-B-C	Warehouse, Room 2100, Insulation From Pipe	None
2100-0282-B-C	Warehouse, Room 2100, Wall	Chrysotile 25
2109-1303-B-C	Warehouse, Room 2109, Insln Piping, Shop Area	Amosite 4
		None
<u>LAB</u>		
1205-0070-B-C	Lab Room, 1205, Pile on Floor, Utility Room	Chrysotile 6
1205-0071-B-C	Lab Room, 1205, Ductwork Insulation	Chrysotile TR <1
1205-0072-B-C	Lab Room, 1205, Around Furnace, Utility Room	Chrysotile 26
1205-0094-B-C	Lab Room, 1205, A/C Filter duct, Inlet	None

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TABLE 3.3-3
SUMMARY OF 2,3,7,8-TCDD RESULTS
BUILDINGS AND STRUCTURES

LOCATION	NUMBER OF SAMPLES	NUMBER OF POSITIVE ANALYSES	RANGE OF CONCENTRATION (ppb)	CHIPS			BULK		
				NUMBER OF SAMPLES	NUMBER OF POSITIVE ANALYSES	RANGE OF CONCENTRATION (ppb)	NUMBER OF SAMPLES	NUMBER OF POSITIVE ANALYSES	RANGE OF CONCENTRATION (ppb)
Office and Laboratory	24(1)	22	10-16,000	16	10	0.57-69.3	-	-	-
Warehouse	8	8	13-19,000	16	13	1.0-192	-	-	-
Manufacturing Building	5	4	233-7,000	23	23	0.93-1,300	-	-	-
Process Building	14	14	6.4-41,600	10	10	2.7-1,500	5	5	3.0-128
Other Structures	-	-	-	6	6	1.2-50.0	1	1	0.17
TOTAL	51	48	6.4-41,600	71	62	0.57-1,500	6	6	0.17-128

(1) One sample void

TABLE 5.3-4
2,3,7,8-TCDD SAMPLE REANALYSIS SUMMARY
BUILDINGS AND STRUCTURES

SAMPLE IDENTIFICATION	LOCATION	SAMPLE TYPE	INITIAL RESULTS (ppb)/ng/m ²	ACTION	REANALYSIS RESULTS (ppb)/ng/m ²
2109-0178-W-L	Warehouse	Wipe	25,000	10:1 dilution	19,000
4504-0452-C-L	Process	Chip	(>938)	1 gram	(1,580)
4100-0553-C-L	Process	Chip	(1,678)	1 gram	(696)
4100-0554-C-L	Process	Chip	(529)	1 gram	(445)
4100-0560-W-L	Process	Wipe	137,000 ^(a)	100:1 dilution	29,200
4100-0560-W-L	Process	Wipe	38,900	10:1 dilution	41,600
3100-0621-C-L	Manufacturing	Chip	(1,316)	1 gram	(1,280)
3100-0633-C-L	Manufacturing	Chip	(482)	1 gram	(447)
3100-0634-C-L	Manufacturing	Chip	(b)	1 gram	(502)
3200-0634-C-L	Manufacturing	Chip	(1,000)	1 gram	(896)

(a) Extremely high matrix interference and saturation.

(b) Instrument detector saturated - no results available.

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TABLE 5.3.1-1
2,3,7,8-TCDD SAMPLE ANALYSIS
OFFICE AND LABORATORY BUILDING

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>WIPE SAMPLES</u>		
1100-0016-W-L	Lab Room 1100, Main Entrance	76 ng/m ²
1102-0017-W-L	Lab Room 1102, Accounting	38 ng/m ²
1105-0018-W-L	Lab Room 1105, Floor, Plant Manager	100 ng/m ²
1106-0021-W-L	Lab Room 1106, Floor, Back Foyer Inside Door	500 ng/m ²
1107-0020-W-L	Lab Room 1107, Floor	100 ng/m ²
1108-0019-W-L	Lab Room 1108, Wall	480 ng/m ²
1116-0034-W-L	Lab Room 1116, Locker Room	500 ng/m ²
1122-0035-W-L	Lab Room 1122, Heater Duct, Basket Room	120 ng/m ²
1122-0073-W-L	Lab Room 1122, Windowsill, Basket Room	520 ng/m ²
1122-0074-W-L	Lab Room 1122, Floor Near Inside Entrance	1,100 ng/m ²
1202-0032-W-L	Lab Room 1202, Floor, Lunchroom	56 ng/m ²
1202-0033-W-L	Lab Room 1202, Radiator, Lunchroom	18 ng/m ²
1204-0023-W-L	Lab Room 1204, Floor by Back Door, Lab	150 ng/m ²
1204-0024-W-L	Lab Room 1204, Lab Hood, Lab	14,000 ng/m ²
1204-0025-W-L	Lab Room 1204, North Side of Entrance, Lab Side	10 ng/m ²
1204-0026-W-L	Lab Room 1204, Bench Near Back Door	1,000 ng/m ²
1205-0030-W-L	Lab room 1205, A/C Intake Duct, Utility	1,200 ng/m ²
1205-0031-W-L	Lab Room 1205, Furnace Intake, Utility Room	88 ng/m ²
1205-0095-W-L	Lab Room 1205, Heater Interior Inlet, Utility	1,400 ng/m ²
1206-0027-W-L	Lab Room 1206, Floor, Small Lab	350 ng/m ²
1206-0028-W-L	Lab Room 1206, Bench, Small Lab	Void sample lost
1206-0381-W-L	Office/Lab Room 1206, Bench, Small Lab	150 ng/m ²
1306-1590-W-L	Office Lab, West Wall, at Roof	168 ng/m ²
1306-0101-W-L	Office Lab, Center of West Wall, at Roof	ND (3.2 ng/m ²)

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TABLE 5.3.1-1
(Continued)

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>CHIP SAMPLES</u>		
1118-0049-C-L	Lab Room 1118, Floor Under Sink Edge, Washroom	2.0 ppb
1119-0050-C-L	Lab room 1119, Floor Slop Sink	3.7 ppb
1122-0051-C-L	Lab Room 1122, Floor Under Arch Between Room 1122 and 1116	25.0 ppb
1122-0052-C-L	Lab Room 1122, Floor Near Drain, Basket Room	69.3 ppb
1122-0053-C-L	Lab Room 1122, Floor Near Back Door, Basket Room	61.2 ppb
1501-0098-C-L	Lab Exterior, 1501, Center, North Wall at Roofsill	ND (0.10 ppb)
1501-0111-C-L	Lab Exterior, 1501, Center of North Wall, 3 to 5 feet	0.70 ppb
1501-0112-C-L	ITAS Split of 1501-0111-C-L	0.95 ppb
1501-0113-C-L	Lab Exterior, 1501, Center North Wall, Ground Level	0.57 ppb
1505-0097-C-L	Lab Exterior, 1505, South Corner, East Wall at Roofsill	ND (0.08 ppb)
1505-0108-C-L	Lab Exterior, 1505, South Corner, East Wall, 3 to 5 feet	ND (0.63 ppb)
1505-0109-C-L	Lab Exterior, 1505, South Corner, East Wall, Ground Level	ND (0.25 ppb)
1505-0110-C-L	Lab Exterior, 1505, Walkway of Front Entrance	2.3 ppb
1506-0099-C-L	Lab Exterior, 1506, Center West Wall, Top 24-inch Vertical	ND (0.34 ppb)
1506-0166-C-L	Lab Exterior, 1506, Center West Wall, 3 to 5 feet	ND (0.58 ppb)
1506-0167-C-L	Lab Exterior, 1506, Center West Wall, Ground Level	2.4 ppb

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TABLE 5.3.2-1
2,3,7,8-TCDD SAMPLE ANALYSIS
WAREHOUSE

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>WIPE SAMPLES</u>		
2100-0218-W-L	Warehouse, Room 2100, Top of Fluorescent	8,120 ng/m ²
2103-0217-W-L	Warehouse, Room 2103, Floor, Foreman's Office	1,810 ng/m ²
2108-0176-W-L	Warehouse, Room 2108, Floor, Kitchen	600 ng/m ²
2108-0177-W-L	Warehouse, Room 2108, Windowsill, Kitchen	130 ng/m ²
2109-0178-W-L	Warehouse, Room 2109, Top of Light Work Area, Shop	19,000 ng/m ²
2109-0179-W-L	Warehouse, Room 2109, Top of Bench in Shop	3,500 ng/m ²
2200-0180-W-L	Warehouse, Room 2200, Top of Beam in Storage Area	8,000 ng/m ²
2400-0315-W-L	Warehouse, West End, Roof	13 ng/m ²
<u>CHIP SAMPLES</u>		
2100-0168-C-L	Warehouse, Room 2100, Center of Traffic Area, Floor	54.6 ppb
2109-0169-C-L	Warehouse, Room 2109, Floor, Tool Crib Cage Area	48.7 ppb
2109-0170-C-L	Warehouse, Room 2109, Floor By Traffic Door	121 ppb
2109-0171-C-L	Warehouse, Room 2109, Floor by Warehouse Door	192 ppb
2501-0317-C-L	Warehouse, North Wall at Ground Level	4.4 ppb
2501-0391-C-L	Warehouse North Wall at 60 inches (3 to 5 feet)	1.6 ppb
2501-0392-C-L	Warehouse Exterior, North Side at Roof	1.9 ppb
2502-0319-C-L	Warehouse, South Wall at Ground Level	10 ppb
2502-0393-C-L	Warehouse, South Wall at 60 inches (3 to 5 feet)	13.3 ppb
2502-0529-C-L	Warehouse, Exterior, South Wall, at Roof Line	16.5 ppb
2504-0318-C-L	Warehouse, East Wall at Ground Level	3.1 ppb

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TABLE 5.3.2-1
(Continued)

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>CHIP SAMPLES</u> (Continued)		
2504-0527-C-L	Warehouse, Exterior, East Wall, 3 to 5 feet	1.4 ppb
2504-0528-C-L	Warehouse, Exterior, East Wall, at Roof Line	1.0 ppb
2506-0316-C-L	West Wall at Ground Level	ND(0.57 ppb)
2506-0389-C-L	Warehouse, West Wall at 60 inches (3 to 5 feet)	ND (0.77 ppb)
2506-0390-C-L	Warehouse, West Wall at Roof Line	ND (0.28 ppb)

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TABLE 5.3.3-1
2,3,7,8-TCDD SAMPLE ANALYSIS
MANUFACTURING BUILDING

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>WIPE SAMPLES</u>		
3100-0658-W-L	Mftg. Bldg., First Floor, Packing Area, Rafter	233 ng/m ²
3200-0655-W-L	Mftg. Bldg., Second Floor, New Addition, Floor, South End	7,000 ng/m ²
3200-0656-W-L	Mftg. Bldg., Second Floor, New Addition, Panel, Center	1,100 ng/m ²
3200-0657-W-L	Mftg. Bldg., Second Floor, New Addition, North End, Beam	630 ng/m ²
3502-0716-W-L	Mftg. Bldg., South Exterior Door	ND (77.5 ng/m ²)
<u>CHIP SAMPLES</u>		
3100-0619-C-L	Mftg. Bldg., Old Area, Roof Slab, South of Center Vessel	1.1 ppb
3100-0620-C-L	Mftg. Bldg., Old Area, Roof Slab, West of North Vessel	12.3 ppb
3100-0621-C-L	Mftg. Bldg., Bulk Debris from Drain Area	1,280 ppb
3100-0622-C-L	Mftg. Bldg., Old Area, 1st Floor, Floor North End North Room	91.8 ppb
3100-0633-C-L	Mftg. Bldg., Old Area, Floor, Center	447 ppb
3100-0634-C-L	Mftg. Bldg., Old Area, Floor, South	502 ppb
3100-0635-C-L	Mftg. Bldg., Packing Area, Floor at Main Door	210 ppb
3100-0636-C-L	Mftg. Bldg., Packing Area, Floor at Packing Chute	191 ppb
3100-0639-C-L	Mftg. Bldg., Packing Area, Low on East Wall	6.0 ppb
3100-0640-C-L	Mftg. Bldg., Packing Area, 30 to 60 inches on West Wall	18.1 ppb
3100-0641-C-L	Mftg. Bldg., New Addition, Southwest Wall, Interior	62.1 ppb
3100-0652-C-L	Mftg. Bldg., First Floor, Southwest Floor Under Vessel	5.1 ppb
3100-0653-C-L	Mftg. Bldg., First Floor, New Addition, Center Floor by Pump	22.5 ppb
3200-0654-C-L	Mftg. Bldg., Second Floor, New Addition, North Wall by Door	896 ppb

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TABLE 5.3.3-1
(Continued)

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>CHIP SAMPLES</u> (Continued)		
3501-0690-C-L	Mftg. Bldg., Exterior, North Wall, 0 to 24 inches by Main Door	203 ppb
3501-0691-C-L	Mftg. Bldg., Exterior, North Wall, 36 to 60 inches by Main Door	167 ppb
3502-0696-C-L	Mftg. Bldg., Exterior, South, Under Load-Out Door	200 ppb
3502-0697-C-L	Mftg. Bldg., Exterior, South Wall, 0 to 24 inches, Package Area Door	6.9 ppb
3502-0698-C-L	Mftg. Bldg., Exterior, South Wall, 36 to 60 inches, Package Area Door	26.6 ppb
3506-0692-C-L	Mftg. Bldg., Exterior, West Wall, 0 to 24 inches, by Large North Door	59.8 ppb
3506-0693-C-L	Mftg. Bldg., Exterior, West Wall, 36 to 60 inches, by Large North Doorway	12.2 ppb
3506-0694-C-L	Mftg. Bldg., Exterior, West Wall, 0 to 24 inches, by South Stairway	3.1 ppb
3506-0695-C-L	Mftg. Bldg., Exterior, West Wall, 36 to 60 inches, by South Stairway	0.93 ppb

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TABLE 5.3.4-1
2,3,7,8-TCDD SAMPLE ANALYSIS
PROCESS BUILDING

SAMPLE NUMBER	DESCRIPTION	RESULTS
WIPE SAMPLES		
4100-0556-W-L	Process Bldg., East End, Near Vessel, Top of Light, First Floor	1,970 ng/m ²
4100-0557-W-L	Process Bldg., East End, Low on Column, Near Vessel, First Floor	4,040 ng/m ²
4100-0558-W-L	Process Bldg., Center, First Floor, Top of Light, Near Vessel	1,200 ng/m ²
4100-0559-W-L	Process Bldg., Center, First Floor, Low on Column, Near Vessel	29,200 ng/m ²
4100-0560-W-L	Process Bldg., West End, First Floor, Top of Light, Near Vessel	41,600 ng/m ²
4100-0561-W-L	Process Bldg., West End, First Floor, Low on Column, Near Vessel	9,070 ng/m ²
4200-0608-W-L	Process Bldg., Second Floor, West End Interior Wall	1,200 ng/m ²
4200-0609-W-L	Process Bldg., Second Floor, Acid Room Wall (Interior)	380 ng/m ²
4200-0610-W-L	Process Bldg., Second Floor, East End Interior Wall	270 ng/m ²
4300-0611-W-L	Process Bldg., Third Floor, East End Interior Wall	3,100 ng/m ²
4300-0612-W-L	Process Bldg., Third Floor, Surface, Center	170 ng/m ²
4300-0613-W-L	Process Bldg., Third Floor, Surface, East End	60 ng/m ²
4400-0496-W-L	Process Bldg., Roof, Northeast Quadrant	6.4 ng/m ²
4400-0496-W-L	Process Bldg., Roof, Southwest Corner	12 ng/m ²
CHIP SAMPLES		
4100-0553-C-L	Process Bldg., Floor, West End of First Floor	696 ppb
4100-0554-C-L	Process Bldg., Floor at Loading Door, First Floor	445 ppb
4100-0555-C-L	Process Bldg., Floor, East End Under Vessel, First Floor	43.2 ppb
4501-0424-C-L	Process Bldg., Exterior, North Wall, 0 to 24 inches	45 ppb
4501-0428-C-L	Process Bldg., In Bin, North, 0 to 24 inches	37.0 ppb

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SAMPLE NUMBER	DESCRIPTION	RESULTS
CHIP SAMPLES (Continued)		
4502-0451-C-L	Process Bldg., South Wall, Near Roof at Vert Stairs	76.8 ppb
4503-0427-C-L	Process Bldg., Exterior, South at C Filler, 24 inches Over Curb	67.9 ppb
4504-0452-C-L	Process Bldg., East Wall, Over Trench Near Vessels (0 to 24 inches)	1,580 ppb
4506-0425-C-L	Process Bldg., Exterior Bin Wall, West Side, 0 to 24 inches	2.7 ppb
4506-0426-C-L	Process Bldg., Exterior, Bin Wall, West Side, 36 to 60 inches	2.9 ppb
BULK SAMPLES		
4501-0455-B-L	Process Bldg., North Wall, 36 to 60 inches	128 ppb
4501-0493-B-L	Process Bldg., North Wall, 24 inches from top (Off Louvers)	3.0 ppb
4503-0456-B-L	Process Bldg., South Wall, 36 to 60 inches	8.1 ppb
4504-0453-B-L	Process Bldg., East Wall, Near Vessels (36 to 60 inches)	95.4 ppb
4504-0454-B-L	Process Bldg., East Wall, at Roof Near Vessels	78.3 ppb

TABLE 5.3.4-1
(Continued)

TABLE 5.3.5-1
2,3,7,8-TCDD RESULTS
OTHER STRUCTURES

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>CHIP SAMPLES</u>		
5001-0277-C-L	Stack Flue, Soot at Furnace Entrance	10.5 ppb
5002-0278-C-L	Stack, Soot from Base of Inside Dropout Chamber	9.2 ppb
5003-0279-C-L	Stack, Exterior at Base, 0 to 24 inches Vertical	1.2 ppb
6100-0388-C-L	Solvent Shed Interior Floor	9.0 ppb
6200-0618-C-L	Pump House, Interior, Floor	50.0 ppb
6600-0617-C-L	Pump House, Exterior, 0 to 24 inches	5.3 ppb
<u>BULK SAMPLES</u>		
6500-0280-B-L	Solvent Shed Exterior, Insulating Panel	0.17 ppb

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TABLE 5.3.6-1
2,3,7,8-TCDD
ANALYSIS RESULTS OF TANK SAMPLES

SAMPLE NUMBER	DESCRIPTION	RESULTS (ppb)
7037-1206-N-L	Tank No. 37, Bulk, reddish-brown sediment	No recovery ⁽¹⁾
7041-1192-N-L	Tank No. 41, Bulk, pink-hardened clay	No recovery ⁽¹⁾
7057-1258-N-L	Tank No. 57, Bulk, whitish-grey solid	5.0
7063-1264-N-L	Tank No. 63, Bulk, white powder	100
7094-1410-N-L	Tank No. 94, Bulk, greyish-green solid	236
7112-1523-N-L	Tank No. 112, Bulk, orange liquid	No recovery ⁽²⁾
7118-1526-N-L	Tank No. 118, Wipe, glass lined	8.2
7126-1539-N-L	Tank No. 126, Bulk, rusty-brown solid	5530
7127-1540-N-L	Tank No. 127, Bulk, red crystals	4200
7129-1548-N-L	Tank No. 129, Bulk, brown, black-rusty solid	679
7135-1620-N-L	Tank No. 135, Bulk, rusty-brown powder	60,800
7136-1635-N-L	Tank No. 136, Bulk, red-rusty solid	11.1

(1) Three analyses attempted no known method available.

(2) No sample available for reanalysis.

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TABLE 5.3.6-2
SUMMARY OF 2,3,7,8-TCDD
REANALYSIS RESULTS

SAMPLE IDENTIFICATION	INITIAL RESULTS	ACTION	REANALYSIS RESULTS
7126-1539-W-L	6,570	1 gram 5:1	5,530
7127-1540-W-L	6,060	1 gram 5:1	4,200
7129-1548-W-L	560	1 gram	679
7135-1620-W-L	>106,000	1 gram 20:1	60,800

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TABLE 5.4-1
SEWERS AND SUMPS
2,3,7,8-TCDD RESULTS SUMMARY

ANALYSIS LOCATION	NUMBER OF SAMPLES	NUMBER OF POSITIVE RESULTS	RANGE OF CONCENTRATION (ppb)
Sewers	4	4	195-4,040
Sumps			
Manufacturing Building	5	5	105-2,950
Process Building	3	3	350-9,160
TOTAL	12	12	19.5-9,160

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TABLE 3.4-2
2,3,7,8-TCDD RESULTS
SEWERS AND SUMPS

SAMPLE NUMBER	DESCRIPTION	RESULTS
<u>SEWER SAMPLES</u>		
8007-1256-Z-L	12 Feet South of Southwest Corner of Manufacturing Building	19.5 ppb
8010-1286-Z-L	25 Feet North of 15 Feet Southwest Outside Wall at Manufacturing Building	4,040 ppb
8011-1287-Z-L	Directly 20 Feet South of Tank No. 23 Near Warehouse	420 ppb
8012-1323-Z-L	50 Feet Northeast of Office Lab	529 ppb
<u>SUMP SAMPLES</u>		
8001-1231-Z-L	Mftg. Bldg., First Floor, West Wall Next to Rollup Door	2,950 ppb
8002-1232-Z-L	Mftg. Bldg., First Floor, West Side, North of Rollup Door	1,010 ppb
8003-1233-Z-L	Mftg. Bldg., First Floor, Southeast Side, Floor Sump, North Sliding Doors	105 ppb
8004-1234-Z-L	Outside Process Building, East Wall, Floor Sump	350 ppb
8005-1254-Z-L	Outside Wall of Process Building, 30 Feet West of Tank 2099	2,680 ppb
8006-1255-Z-L	Outside Northwest Corner Process Bldg., Five Feet East of Back Stairway	9,160 ppb
8008-1284-Z-L	15 Feet Northwest of Southwest Corner of Manufacturing Building	560 ppb
8009-1285-Z-L	60 Feet North of Outside Southwest Corner of Manufacturing Building	836 ppb

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TABLE 5.4-3
SEWER AND SUMP SAMPLE
2,3,7,8-TCDD
REANALYSIS RESULTS SUMMARY

SAMPLE NUMBER	INITIAL RESULTS (ppb)	ACTION	REANALYSIS RESULTS (ppb)
8001-1231-Z-L	>1,140	1 gram	2,950
8002-1232-Z-L	>1,623	1 gram	1,010
8004-1234-Z-L	>361	1 gram	350
8005-1254-Z-L	>2,302	1 gram	2,680
8006-1255-Z-L	>5,530	Dilution 50:1	9,160
8008-1284-Z-L	>1,280	1 gram	560
8010-1286-Z-L	>3,660	Dilution	4,040
8011-1287-Z-L	>567	1 gram	420
8012-1323-Z-L	>386	1 gram	529

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TABLE 5.5.2-1
DESIGNATED ANALYSIS FOR NEAR-SURFACE AND BORING SOIL SAMPLES

SAMPLE ELEVATION CODE	RELATIVE SAMPLE DEPTH (inches)	ANALYSES DESIGNATED	
		NSSS	BSS
100	0-6	Priority Pollutants ²	Priority Pollutants
101	6-12	Dioxin ²	Dioxin
102	12-24	Priority Pollutants	Priority Pollutants
103-108		(24-60") Samples archived	(24"-last increment above silt) Samples archived
109	above silt zone	-	Total
201	silt zone	-	Dioxin

(1) Priority pollutant refers to the analysis and compounds contained therein as described by priority pollutants (HSL compound) for acid/base/neutral; volatile organic compounds, pesticides and PCB, metals, total cyanide, total phenols, herbicides (8150), 2,3,7,8-tetrachlorodibenzo-p-dioxin.

(2) Dioxin refers to the analysis for 2,3,7,8-tetrachlorodibenzo-p-dioxin.

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TABLE 5.5.2.1-1
COLLECTED NEAR-SURFACE SOIL SAMPLES

DEPTH (inches)	ELEVATION CODE	NUMBER OF SAMPLES
0-6	100	21
6-12	101	21
12-24	102	21
24-36	103	21
36-48	104	17
48-60	105	<u>14</u>
TOTAL		115

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TABLE 5.5.2.1-2
NEAR-SURFACE SOIL SAMPLE 2,3,7,8-TCDD REANALYSIS SUMMARY

STATION NUMBER	DEPTH (inches)	ELEVATION CODE	INITIAL RESULTS (ppb)	CORRECTIVE ACTION	REANALYSIS RESULTS (ppb)
A-2-G	0-6	100	296	1 gram	326
A-2-G	6-12	101	289	1 gram	330
A-5-G	0-6	100	500	1 gram	695
A-5-G	6-12	101	460	1 gram	453
F-5-E	0-6	100	268	1 gram	470
F-5-E	6-12	101	247	1 gram	394
F-5-E	12-24	102	>19,000	1 gram, dilution 10:1	19,500
G-3-I	0-6	100	1,110	1 gram	1,010
G-3-L	0-6	100	261	1 gram	310
G-4-A	0-6	100	395	1 gram	276
G-4-A	6-12	101	>3,130	1 gram, dilution 3:1	3,690
G-4-A	12-24	102	>1,515	1 gram	1,770
G-5-P	0-6	100	325	1 gram	361
G-5-P	6-12	101	359	1 gram	494
H-2-H	0-6	100	>1,586	1 gram	2,390
H-2-H	6-12	101	1,180	1 gram	1,230
H-2-H	12-24	102	286	1 gram	510
H-5-F	12-24	102	336	1 gram	385
H-7-P	0-6	100	>5,768	1 gram, dilution 5:1	9,050
H-7-P	6-12	101	>1,550	1 gram	2,730
H-7-P	12-24	102	231	1 gram	200

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A - The sample was analyzed for possible future analysis.
B - Refused, then a sample was not recovered.

SAMPLE CODE	ELEVATION (feet)	DEPTH (inches)	100	101	102	103	104	105
A-2-G	120	0.39	695	163	153	153	153	153
A-4-F	120	0.39	695	163	153	153	153	153
A-5-G	120	0.39	695	163	153	153	153	153
B-2-H	120	0.39	695	163	153	153	153	153
C-2-B	120	0.39	695	163	153	153	153	153
D-4-H	120	0.39	695	163	153	153	153	153
E-1-G	120	0.39	695	163	153	153	153	153
F-5-B	120	0.39	695	163	153	153	153	153
G-3-I	120	0.39	695	163	153	153	153	153
H-1-L	120	0.39	695	163	153	153	153	153
I-4-A	120	0.39	695	163	153	153	153	153
J-5-E	120	0.39	695	163	153	153	153	153
K-3-F	120	0.39	695	163	153	153	153	153
L-1-N	120	0.39	695	163	153	153	153	153
M-2-A	120	0.39	695	163	153	153	153	153
N-2-B	120	0.39	695	163	153	153	153	153
O-5-F	120	0.39	695	163	153	153	153	153
P-2-H	120	0.39	695	163	153	153	153	153
Q-3-E	120	0.39	695	163	153	153	153	153
R-4-A	120	0.39	695	163	153	153	153	153
S-5-E	120	0.39	695	163	153	153	153	153
T-3-L	120	0.39	695	163	153	153	153	153
U-4-A	120	0.39	695	163	153	153	153	153
V-5-E	120	0.39	695	163	153	153	153	153
W-3-F	120	0.39	695	163	153	153	153	153
X-1-N	120	0.39	695	163	153	153	153	153
Y-2-A	120	0.39	695	163	153	153	153	153
Z-2-B	120	0.39	695	163	153	153	153	153
AA-5-F	120	0.39	695	163	153	153	153	153
AB-2-H	120	0.39	695	163	153	153	153	153
AC-3-E	120	0.39	695	163	153	153	153	153
AD-4-A	120	0.39	695	163	153	153	153	153
AE-5-E	120	0.39	695	163	153	153	153	153
AF-3-L	120	0.39	695	163	153	153	153	153
AG-4-A	120	0.39	695	163	153	153	153	153
AH-5-E	120	0.39	695	163	153	153	153	153
AI-3-F	120	0.39	695	163	153	153	153	153
AJ-1-N	120	0.39	695	163	153	153	153	153
AK-2-A	120	0.39	695	163	153	153	153	153
AL-2-B	120	0.39	695	163	153	153	153	153
AM-5-F	120	0.39	695	163	153	153	153	153
AN-2-H	120	0.39	695	163	153	153	153	153
AO-3-E	120	0.39	695	163	153	153	153	153
AP-4-A	120	0.39	695	163	153	153	153	153
AQ-5-E	120	0.39	695	163	153	153	153	153
AR-3-L	120	0.39	695	163	153	153	153	153
AS-4-A	120	0.39	695	163	153	153	153	153
AT-5-E	120	0.39	695	163	153	153	153	153
AU-3-F	120	0.39	695	163	153	153	153	153
AV-1-N	120	0.39	695	163	153	153	153	153
AW-2-A	120	0.39	695	163	153	153	153	153
AX-2-B	120	0.39	695	163	153	153	153	153
AY-5-F	120	0.39	695	163	153	153	153	153
AZ-2-H	120	0.39	695	163	153	153	153	153
BA-3-E	120	0.39	695	163	153	153	153	153
BB-4-A	120	0.39	695	163	153	153	153	153
BC-5-E	120	0.39	695	163	153	153	153	153
BD-3-L	120	0.39	695	163	153	153	153	153
BE-4-A	120	0.39	695	163	153	153	153	153
BF-5-E	120	0.39	695	163	153	153	153	153
BG-3-F	120	0.39	695	163	153	153	153	153
BH-1-N	120	0.39	695	163	153	153	153	153
BI-2-A	120	0.39	695	163	153	153	153	153
BJ-2-B	120	0.39	695	163	153	153	153	153
BK-5-F	120	0.39	695	163	153	153	153	153
BL-2-H	120	0.39	695	163	153	153	153	153
BM-3-E	120	0.39	695	163	153	153	153	153
BN-4-A	120	0.39	695	163	153	153	153	153
BO-5-E	120	0.39	695	163	153	153	153	153
BP-3-L	120	0.39	695	163	153	153	153	153
BQ-4-A	120	0.39	695	163	153	153	153	153
BR-5-E	120	0.39	695	163	153	153	153	153
BS-3-F	120	0.39	695	163	153	153	153	153
BT-1-N	120	0.39	695	163	153	153	153	153
BU-2-A	120	0.39	695	163	153	153	153	153
BV-2-B	120	0.39	695	163	153	153	153	153
BW-5-F	120	0.39	695	163	153	153	153	153
BX-2-H	120	0.39	695	163	153	153	153	153
BY-3-E	120	0.39	695	163	153	153	153	153
BZ-4-A	120	0.39	695	163	153	153	153	153
CA-5-E	120	0.39	695	163	153	153	153	153
CB-3-L	120	0.39	695	163	153	153	153	153
CC-4-A	120	0.39	695	163	153	153	153	153
CD-5-E	120	0.39	695	163	153	153	153	153
CE-3-F	120	0.39	695	163	153	153	153	153
CF-1-N	120	0.39	695	163	153	153	153	153
CG-2-A	120	0.39	695	163	153	153	153	153
CH-2-B	120	0.39	695	163	153	153	153	153
CI-5-F	120	0.39	695	163	153	153	153	153
CJ-2-H	120	0.39	695	163	153	153	153	153
CK-3-E	120	0.39	695	163	153	153	153	153
CL-4-A	120	0.39	695	163	153	153	153	153
CM-5-E	120	0.39	695	163	153	153	153	153
CN-3-L	120	0.39	695	163	153	153	153	153
CO-4-A	120	0.39	695	163	153	153	153	153
CP-5-E	120	0.39	695	163	153	153	153	153
CQ-3-F	120	0.39	695	163	153	153	153	153
CR-1-N	120	0.39	695	163	153	153	153	153
CS-2-A	120	0.39	695	163	153	153	153	153
CT-2-B	120	0.39	695	163	153	153	153	153
CU-5-F	120	0.39	695	163	153	153	153	153
CV-2-H	120	0.39	695	163	153	153	153	153
CW-3-E	120	0.39	695	163	153	153	153	153
CX-4-A	120	0.39	695	163	153	153	153	153
CY-5-E	120	0.39	695	163	153	153	153	153
CZ-3-L	120	0.39	695	163	153	153	153	153
DA-4-A	120	0.39	695	163	153	153	153	153
DB-5-E	120	0.39	695	163	153	153	153	153
DC-3-F	120	0.39	695	163	153	153	153	153
DD-1-N	120	0.39	695	163	153	153	153	153
DE-2-A	120	0.39	695	163	153	153	153	153
DF-2-B	120	0.39	695	163	153	153	153	153
DG-5-F	120	0.39	695	163	153	153	153	153
DH-2-H	120	0.39	695	163	153	153	153	153
DI-3-E	120	0.39	695	163	153	153	153	153
DJ-4-A	120	0.39	695	163	153	153	153	153
DK-5-E	120	0.39	695	163	153	153	153	153
DL-3-L	120	0.39	695	163	153	153	153	153
DM-4-A	120	0.39	695	163	153	153	153	153
DN-5-E	120	0.39	695	163	153	153	153	153
DO-3-F	120	0.39	695	163	153	153	153	153
DP-1-N	120	0.39	695	163	153	153	153	153
DQ-2-A	120	0.39	695	163	153	153	153	153
DR-2-B	120	0.39	695	163	153	153	153	153
DS-5-F	120	0.39	695	163	153	153	153	153
DT-2-H	120	0.39	695	163	153	153	153	153
DU-3-E	120	0.39	695	163	153	153	153	153
DV-4-A	120	0.39	695	163	153	153	153	153
DW-5-E	120	0.39	695	163	153	153	153	153
DX-3-L	120	0.39	695	163	153	153	153	153
DY-4-A	120	0.39	695	163	153	153	153	153
DZ-5-E	120	0.39	695	163	153	153	153	153
EA-3-F	120	0.39	695	163	153	153	153	153
EB-1-N	120	0.39	695	163	153	153	153	153
EC-2-A	120	0.39	695	163	153	153	153	153
ED-2-B	120	0.39	695	163	153	153	153	153
EE-5-F	120	0.39	695	163	153	153	153	153
EF-2-H	120	0.39	695	163	153	153	153	153
EG-3-E	120	0.39	695	163	153	153	153	153
EH-4-A	120	0.39	695	163	153	153	153	153
EI-5-E	120	0.39	695	163	153	153	153	153
EJ-3-L	120	0.39	695	163	153	153	153	153
EK-4-A	120	0.39	695	163	153	153	153	153
EL-5-E	120	0.39	695	163	153	153	153	153
EM-3-F	120	0.39	695	163	153	153	153	153
EN-1-N	120	0.39	695	163	153	153	153	153
EO-2-A	120	0.39	695	163	153	153	153	153
EP-2-B	120	0.39	695	163	153	153	153	153
EQ-5-F	120	0.39	695	163	153	153	153	153
ER-2-H	120	0.39	695	163	153	153	153	153
ES-3-E	120	0.39	695	163	153	153	153	153
ET-4-A								

TABLE 5.5.2.1-4
NEAR-SURFACE SOILS
ORGANIC PRIORITY POLLUTANT ANALYSIS LEVELS

STATION	DEPTH (inches)	VOA	BASE/NEUTRAL/ACID	PESTICIDE	HERBICIDE
A-2-G	0-6	Low	Medium	Medium ⁽¹⁾	Low ⁽²⁾
A-2-G	12-24	Low	Medium	Medium ⁽³⁾	Low ⁽²⁾
A-4-F	0-6	Low	Low	Low ⁽⁷⁾	Low
A-4-F	12-24	Low	Medium	Medium ⁽⁷⁾	Low
A-5G	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
A-5G	12-24	Low	Medium	Medium	Low
B-2-M	0-6	Low	Medium	Medium ⁽⁴⁾	Low ⁽²⁾
B-2-M	12-24	Low	Low	Low ⁽⁵⁾	Low ⁽²⁾
C-6-B	0-6	Low	Low	Low	Low
C-6-B	12-24	Medium ⁽⁴⁾	Medium	Medium	Low ⁽⁵⁾
D-4-M	0-6	Low	Low	Low ⁽⁷⁾	Low
D-4-M	12-24	Low	Low	Low ⁽⁷⁾	Low
E-1-G	0-6	Low	Low	Low ⁽⁵⁾	Low
E-1-G	12-24	Low	Medium	Medium ⁽⁴⁾	Low
E-5-D	0-6	Low	Medium	Medium ⁽⁷⁾	Low ⁽²⁾
E-5-D	12-24	Low	Low	Low ⁽⁸⁾	Low ⁽²⁾
F-5-E	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
F-5-E	12-24	Low	Medium	Medium ⁽⁶⁾	Low ⁽³⁾
G-3-I	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
G-3-I	12-24	Low	Low	Low ⁽⁷⁾	Low ⁽²⁾
G-3-L	0-6	Low	Low	Low ⁽¹²⁾	Low ⁽²⁾
G-3-L	12-24	Low	Low	Low ⁽⁹⁾	Low ⁽²⁾
G-4-A	0-6	Low	Low	Low ⁽⁹⁾	Low ⁽²⁾
G-4-A	12-24	Low	Medium	Medium ⁽⁶⁾	Low ⁽²⁾
G-5-E	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
G-5-E	12-24	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
G-5-F	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
G-5-F	12-24	Low	Medium	Medium ⁽⁶⁾	Low ⁽⁹⁾

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TABLE 5.5.2.1-4
(Continued)

STATION	DEPTH (inches)	VOA	BASE/NEUTRAL/ACID	PESTICIDE	HERBICIDE
H-1-H	0-6	Low	Low	Low ⁽⁶⁾	Low ⁽²⁾
H-1-H	12-24	Low	Low	Low ⁽⁴⁾	Low
H-2-B	0-6	Low	Medium	Medium ⁽⁴⁾	Low ⁽⁹⁾
H-2-B	12-24	Low	Medium	Medium ⁽⁹⁾	Low ⁽⁹⁾
H-2-H	0-6	Low	Medium	Medium ⁽⁶⁾	Low ⁽⁶⁾
H-2-H	12-24	Low	Medium	Medium ⁽⁶⁾	Low ⁽¹⁰⁾
H-5-F	0-6	Low	Low	Low ⁽⁶⁾	Low
H-5-F	12-24	Low	Medium	Low ⁽⁷⁾	Low ⁽²⁾
H-7-F	0-6	Medium	Medium ⁽⁴⁾	Medium ⁽¹⁾	Low ⁽¹¹⁾
H-7-F	12-24	Low ⁽⁴⁾	Medium	Medium ⁽⁷⁾	Low ⁽¹⁾
H-7-H	0-6	Low	Low	Low ⁽⁷⁾	Low ⁽²⁾
H-7-H	12-24	Medium	Medium ⁽⁷⁾	Medium ⁽¹⁾	Low ⁽¹⁾
J-6-K	0-6	Low	Low	Low ⁽⁷⁾	Low ⁽²⁾
J-6-K	12-24	Low	Low	Low	Low ⁽²⁾

- (1) Further dilution 1:1000
- (2) Further dilution 1:5
- (3) Further dilution 1:500
- (4) Further dilution 1:20
- (5) Further dilution 1:40
- (6) Further dilution 1:100
- (7) Further dilution 1:10
- (8) Further dilution 1:250
- (9) Further dilution 1:50
- (10) Further dilution 1:25
- (11) Further dilution 1:10,000
- (12) Further dilution 1:5000

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0-6 INCHES				12-24 INCHES			
CONCENTRATION RANGE		NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	CONCENTRATION RANGE		NUMBER POSITIVE RESULTS	ANALYZED SAMPLES
2,4,6-Trichlorophenol	1,500,000-1,300	5	21	1,700,000-8,700	4	21	21
2,4-Dichlorophenol	3,600,000-980	7	21	2,500,000-870	8	21	21
2,4-Dimethylphenol	-	0	21	1,700,000	1	21	21
Benzoic Acid	1,800	1	21	-	0	21	21
2,4,5-Trichlorophenol	15,000,000-870	5	21	7,500,000-2,500	5	21	21
Acenaphthene	250	1	21	-	0	21	21
1,2,4-Trichlorobenzene	17,000-1,500	2	21	19,000	1	21	21
Hexachlorobenzene	110,000-560	13	21	720,000-3,200	9	21	21
1,2-Dichlorobenzene	520-230	2	21	9,000	1	21	21
1,3-Dichlorobenzene	-	0	21	610	1	21	21
1,4-Dichlorobenzene	1,400-470	3	21	1,300	1	21	21
Fluoranthene	6,100-330	5	21	64,000-670	6	21	21
Naphthalene	200	1	21	8,200	1	21	21
Di(2-ethylhexyl)phthalate	1,300-310	3	21	310,000-3,100	3	21	21
Di-n-butylphthalate	-	0	21	370,000-2,000	2	21	21
Benzo(a)anthracene	47,000-910	3	21	47,000-310	3	21	21

TABLE 5.5.2.1-5
SUMMARY OF DETECTED BASE/NEUTRAL/ACID ORGANIC COMPOUNDS
NEAR-SURFACE SOILS
(Expressed as ug/kg or ppb)

TABLE 5.5.2.1-5

(Continued)

	0-6 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	SAMPLES
Benzo(a)pyrene	4,800-1,000	3	21	44,000-560	5	21
Benzo(b)fluoranthene	7,100-2,100	3	21	71,000-940	5	21
Chrysene	12,000-2,600	2	21	120,000-1,400	6	21
Acenaphthylene	690-210	2	21	860-240	2	21
Anthracene	3,000-310	4	21	1,200-630	3	21
Benzo(g,h,i)perylene	11,000-3,300	3	21	32,000	1	21
Fluorene	320	1	21	300-250	2	21
Phenanthrene	4,100-250	5	21	61,000-440	6	21
Indeno(1,2,3,-CD)-pyrene	2,500-2,200	2	21	21,000-480	2	21
Pyrene	2,200-230	6	21	78,000-280	7	21
Dibenzofuran	-	0	21	450	1	21
2-Methylnaphthalene	220	1	21	21,000	1	21

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TABLE 5.5.2.1-6
SUMMARY OF DETECTED VOLATILE ORGANICS
NEAR-SURFACE SOILS
(Expressed as $\mu\text{g/kg}$ or ppb)

	0-6 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Benzene	21	1	21	23,000-11	3	21
Chlorobenzene	84,000-39	2	21	170,000-22	6	21
Chloroform	38	1	21	38,000-13	2	21
Ethylbenzene	-	0	21	60,000	1	21
Methylene chloride	1,500-14	21	21	130,000-21	21	21
Tetrachloroethane	860	1	21	36,000-1,300	2	21
Toluene	-	0	21	2,000,000-7	6	21
Trichloroethene	-	0	21	9	1	21
Acetone	5,000-58	13	21	2,000-68	15	21
2-Butanone	1,400-130	2	21	9,200-51	6	21
Carbon disulfide	-	0	21	7	1	21
2-Hexanone	-	0	21	36,000	1	21
Total xylenes	-	0	21	110,000	1	21

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TABLE 5.5.2.1-7
SUMMARY OF DETECTED HERBICIDES, PESTICIDES, AND PCB's
NEAR-SURFACE SOILS
(Expressed as $\mu\text{g/kg}$ or ppb)

	0-6 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
4,4'-DDT	3,500,000-620	19	21	5,090,000-1,400	15	21
4,4'-DDE	93,000-20	14	21	37,000-1,200	8	21
4,4'-DDD	13,000-1,700	3	21	164,000-1,200	5	21
Alpha-Endosulfan	8,900	1	21	1,400	1	21
Dalapon	70,000-190	9	21	29,000-420	9	21
2,4-D	7,600-740	10	21	85,000-190	13	21
2,4,5-T	2,300-190	9	21	86,000-490	10	21

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TABLE 5.5.2.1-8
SUMMARY OF DETECTED INORGANIC PARAMETERS
NEAR-SURFACE SOILS
(Expressed as ug/kg or ppb)

	0-6 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Antimony	6.6-0.09	14	21	3.0-0.10	17	21
Arsenic	23-0.13	21	21	4-0.60	21	21
Beryllium	0.85-0.22	11	21	0.84-0.25	9	21
Cadmium	3.9-0.09	12	21	26-0.08	14	21
Chromium	50-1.1	21	21	50-5.9	21	21
Copper	260-2.4	21	21	250-2.0	20	21
Lead	887-1.8	21	21	646-2.1	20	21
Mercury	39-0.1	18	21	37-0.4	16	21
Nickel	82-3.1	20	21	40-2.1	20	21
Selenium	48	1	21	2.2-0.01	3	21
Silver	1.2-0.24	7	21	11-0.25	6	21
Zinc	29,000-20	21	21	1,300-8.0	21	21
Total Cyanide	1.97-0.15	19	21	2.8-0.10	19	21
Total Phenols	47.8-0.28	20	21	3,378-0.10	21	21

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TABLE 5.5.2.2-1
SUMMARY OF BORING SOIL SAMPLES
COLLECTED FOR CHEMICAL ANALYSES

DEPTH (inches)	ELEVATION CODE	NUMBER OF SAMPLES
0-6	100	8
6-12	102	8
12-24	103	8
-	109	8
-	201	7
TOTAL		39

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TABLE 5.5.2.2-2
BORING SOIL SAMPLE 2,3,7,8-TCDD REANALYSIS SUMMARY

STATION NUMBER	DEPTH (inches)	ELEVATION CODE	INITIAL RESULTS (ppb)	CORRECTIVE ACTION	REANALYSIS RESULTS (ppb)
C-7-C	6-12	101	477	1 gram	784
F-7-B	0-6	100	>1,450	1 gram	2,560
F-7-B	12-24	102	600	1 gram	687
I-2-L	0-6	100	>1,340	1 gram	2,700
I-5-A	0-6	100	>260	1 gram	523
I-5-A	6-12	101	>530	1 gram	883
I-5-A	12-24	102	>450	1 gram	830
I-7-K	0-6	100	350	1 gram	350
I-7-K	6-12	101	>1,500	1 gram, 3:1 dilution	3,510

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TABLE 3.5.2.2-3
BORING SOIL SAMPLES
2,3,7,8-TCDD RESULTS
(ug/kg or ppb)

SAMPLE ELEVATION CODE	DEPTH (inches)	STATION NUMBERS							
		A-2-E	A-3-C	C-7-C	D-1-F	F-7-B	1-2-L	1-5-A	1-7-K
100	0-6	56.3	19.7	130	61.6	2,560	2,700	523	350
101	6-12	36.0	18.8	784	7.5	109	218	883	3,510
102	12-24	72.5	7.4	247	4.7	687	93.6	830	59.3
109	above silt	0.36 (6.5-8.5')	ND (0.02) (6.5-8.0')	71.8 (6.5-8.0')	0.78 (6.5-8.7')	2.4 (6.5-8')	12.1 (13.5-15.5')	20.9 (13.5-15.2')	5.8 (7-8.5')
201	silt	ND (0.07) (12.7-16.7')	ND (0.3) (11.0-13.0')	2.1 (10.0-12.0')	ND (0.06) (10.7-12.7')	0.49 (10.0-12.0')	2.2 (17.0-19.0')	no sample	2.8 (13.5-15.2')

TABLE 5.5.2.2-4
BORING SOILS SAMPLES
ORGANIC PRIORITY POLLUTANT ANALYSIS LEVELS

STATION	BOREHOLE	DEPTH	VOA	BASE/NEUTRAL/ACID	PESTICIDE	HERBICIDE
I-2-L	1	0-6"	Low	Low	Low ⁽¹⁾	Low ⁽²⁾
I-2-L	1	12-24"	Low	Low	Low ⁽³⁾	Low ⁽²⁾
I-2-L	1	13.5-15.5'	Low ⁽⁴⁾	Low ⁽⁵⁾	Low ⁽³⁾	Low ⁽²⁾
I-5-A	2	0-6"	Low	Low	Low ⁽⁸⁾	Low ⁽²⁾
I-5-A	2	12-24"	Low	Low ⁽⁵⁾	Low ⁽¹¹⁾	Low ⁽²⁾
I-5-A	2	13.5-15.2'	Medium ⁽⁵⁾	Medium ⁽⁵⁾	Medium ⁽⁴⁾	Low ⁽³⁾
I-7-K	3	0-6"	Low	Low ⁽²⁾	Low ⁽¹⁰⁾	Low ⁽⁶⁾
I-7-K	3	12-24"	Low	Low	Low ⁽⁷⁾	Low ⁽⁶⁾
I-7-K	3	7-8.5'	Low	Low	Low ⁽⁵⁾	Low ⁽²⁾
C-7-C	4	0-6"	Low	Low ⁽⁵⁾	Medium	Low
C-7-C	4	12-24"	Low	Low	Medium ⁽⁶⁾	Low
C-7-C	4	6.5-8'	Low ⁽⁷⁾	Low ⁽⁶⁾	Medium ⁽⁶⁾	Low ⁽⁷⁾
A-2-K	5	0-6"	Low	Medium	Medium ⁽¹⁾	Low
A-2-K	5	12-24"	Low	Medium	Medium ⁽⁹⁾	Low
A-2-K	5	6.5-8.5'	Low	Medium	Low ⁽⁶⁾	Low
A-3-C	6	0-6"	Low	Medium	Medium ⁽⁶⁾	Low
A-3-C	6	12-24"	Low	Medium	Medium ⁽⁸⁾	Low
A-3-C	6	6.5-8.5'	Low	Medium	Medium	Low
D-1-F	7	0-6"	Low	Low	Medium	Low
D-1-F	7	12-24"	Low	Medium	Medium	Low
D-1-F	7	6.5-8.7'	Low	Low	Medium	Low
F-7-B	8	0-6"	Low	Low	Low ⁽¹⁰⁾	Low ⁽²⁾
F-7-B	8	12-24"	Low ⁽⁹⁾	Low ⁽⁵⁾	Low ⁽¹⁰⁾	Low ⁽²⁾
F-7-B	8	6.5-8'	Low	Low	Low ⁽⁵⁾	Low ⁽²⁾

- (1) Further dilution 1:500
(2) Further dilution 1:5
(3) Further dilution 1:1000
(4) Further dilution 1:200
(5) Further dilution 1:2
(6) Further dilution 1:10
(7) Further dilution 1:50
(8) Further dilution 1:2000
(9) Further dilution 1:20
(10) Further dilution 1:100
(11) Further dilution 1:10,000

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0-6 INCHES				12-24 INCHES				ABOVE SILT			
NUMBER	POSITIVE	RESULTS	ANALYZED	NUMBER	POSITIVE	RESULTS	ANALYZED	NUMBER	POSITIVE	RESULTS	ANALYZED
CONCENTRATION RANGE				CONCENTRATION RANGE				CONCENTRATION RANGE			
2,4,6-Trichlorophenol	32,000-1,300	3	0	4,400-3,600	2	0	0	360,000-2,000	3	0	0
2-Chlorophenol	2,000-230	2	0	820	1	0	0	6,000-1,200	2	0	0
2,4-Dichlorophenol	98,000-3,900	3	0	27,000-4,700	3	0	0	1,400,000-1,700	5	0	0
Phenol	3,100	1	0	12,000-1,400	2	0	0	13,000-820	2	0	0
2,4,5-Trichlorophenol	20,000-1,500	4	0	16,000-1,600	3	0	0	270,000-12,000	4	0	0
Acenaphthene	2,200	1	0	4,600	1	0	0	0	0	0	0
1,2,4-Trichlorobenzene	1,100-430	2	0	8,500-580	2	0	0	14,000	1	0	0
Hexachlorobenzene	35,000-6,500	5	0	84,000-4,900	4	0	0	30,000	1	0	0
2-Chlorophenyl ether	1,100	1	0	850	1	0	0	-	-	0	0
1,2-Dichlorobenzene	770	1	0	8,600-570	2	0	0	13,000	1	0	0
1,3-Dichlorobenzene	-	0	0	780	1	0	0	3,400	1	0	0
1,4-Dichlorobenzene	2,700	1	0	49,000-960	3	0	0	28,000-4,600	3	0	0
Fluoranthene	8,700-400	5	0	20,000-3,200	4	0	0	1,300-560	3	0	0
Naphthalene	1,300	1	0	11,000	1	0	0	10,000-260	5	0	0
8-(2-ethylhexyl)phthalate	16,000	1	0	5,100-2,600	2	0	0	-	0	0	0

TABLE 3.3.2.2-3
SUMMARY OF DETECTED BASE/NEUTRAL/ACID ORGANICS
IN SOIL BONINGS
(Expressed as mg/kg or ppb)

TABLE S.5.2.2-5
(Continued)

	0-6 INCHES			12-24 INCHES			ABOVE SILT		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Benzo(a)anthracene	-	0	8	1,900	1	8	-	0	8
Benzo(a)pyrene	-	0	8	1,600	1	8	-	0	8
Benzo(b)fluoranthene	-	0	8	7,400	1	8	1,900	1	8
Chrysene	-	0	8	4,200	1	8	-	0	8
Anthracene	950	1	8	1,200	1	8	-	0	8
Fluorene	2,100	1	8	4,200	1	8	-	0	8
Phenanthrene	3,800-230	3	8	14,000-720	5	8	2,200-350	2	8
Indeno(1,2,3,-CD)-pyrene	-	0	8	1,400	1	8	-	0	8
Pyrene	8,100-270	5	8	18,000-1,300	5	8	460-420	2	8
Benzyl Alcohol	-	0	8	20,000	1	8	41,000	1	8
Dibenzofuran	1,300	1	8	2,100	1	8	-	0	8
2-Methylnaphthalene	2,600	1	8	8,000-850	3	8	14,000-1,600	4	8

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SUMMARY OF DETECTED VOLATILE ORGANICS IN SOIL BORINGS				(Expressed as ug/kg or ppb)			
0-6 INCHES				12-24 INCHES			
CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE
Benzene	26	1	1,700-680	22,000-5,600	2	8	2
Chlorobenzene	330	1	26,000-49	100,000-17	4	8	5
Ethylbenzene	-	0	100	14,000-220	1	8	2
Methylchloride	410-38	0	1,600-6	11,000-48	0	8	8
Tetrachloroethane	-	0	15	-	0	8	0
Toluene	12-7	2	2,400-9	180,000-11	4	8	2
Acetone	160-57	5	2,300-110	4,500-85	7	8	6
2-Butanone	-	0	8,900	20,000-6,900	1	8	2
Carbon disulfide	-	0	7	13	1	8	1
Total naphenes	-	0	580	1,200	1	8	8

ABOVE SILT

TABLE 3.3.2.2-7
SUMMARY OF DETECTED HERBICIDES, PESTICIDES, AND PCB'S
IN SOIL BORINGS
(Expressed as ug/kg or ppb)

	0-6 INCHES			12-24 INCHES			ABOVE SILT		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
4,4'-DDT	830,000-17,000	5	8	3,200,000-43,000	5	8	140,000-100	4	8
4,4'-DDE	57,900-6,500	6	8	297,000-2,400	6	8	1,500-290	4	8
4,4'-DDD	78,000-2,000	5	8	182,000-3,900	5	8	370,000-42	5	8
Beta-BHC	130,000-830	2	8	120,000	1	8	100,000	1	8
Delapen	21,000-160	6	8	94,000-300	5	8	-	0	8
Dicamba	1,700-230	3	8	1,600-100	3	8	160	1	8
2,4-D	120,000-240	8	8	16,000-110	8	8	2,600,000-140	7	8
2,4,5-T	54,000-94	8	8	14,000-95	7	8	690,000-610	5	8
2,4-DB	-	0	8	1,400	1	8	170	1	8
Dinoseb (DNBP)	590-210	2	8	-	0	8	-	0	8

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TABLE 5.5.2.2-8
SUMMARY OF DETECTED INORGANIC PARAMETERS
IN SOIL BORINGS
(Expressed as ug/kg or ppb)

	0-6 INCHES			12-24 INCHES			ABOVE SILT		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Antimony	11-0.2	8	8	3.5-0.1	8	8	1.1-0.1	6	8
Arsenic	28-1.0	8	8	26-2.1	8	8	120-5.7	8	8
Beryllium	-	0	8	3.7-0.2	5	8	1.4-0.1	5	8
Cadmium	3-0.5	8	8	2.5-0.3	8	8	3-0.1	6	8
Chromium	72-7.9	8	8	40-13	8	8	25-5.5	8	8
Copper	290-46	8	8	730-82	8	8	6,600-24	8	8
Lead	1,400-73	8	8	2,300-180	8	8	11,000-19	8	8
Mercury	11-0.1	8	8	7.6-0.5	8	8	95-0.2	7	8
Nickel	95-15	8	8	170-13	8	8	72-5.8	8	8
Silver	0.92-0.2	6	8	0.9-0.3	4	8	1.8-0.4	5	8
Zinc	3,900-180	8	8	1,500-190	8	8	1,300-45	8	8
Total Cyanides	1.2-0.25	8	8	3.7-0.15	8	8	1.2-0.1	8	8
Total Phenols	13-0.2	8	8	12-0.2	8	8	1,600-0.3	7	8

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TABLE 5.5.2.3-1
ADDITIONAL SELECTED SILT ZONE SAMPLES FOR
2,3,7,8-TCDD ANALYSIS
(ug/kg or ppb)

SAMPLE LOCATION	BOREHOLE NUMBER	ELEVATION CODE			
		200 ⁽²⁾	201 ⁽¹⁾	202 ⁽²⁾	203 ⁽²⁾
C-7-C	4		2.1 (10.0-12.0')	1.2 (12.0-14.0')	
Initial F-5-E ⁽³⁾	11		1.8 (silt)		
Archive F-5-E			WD (0.24) (10.5-12.5')		WD (0.18) (141.5-16.5')
G-5-E ⁽⁴⁾	10		11.8 (silt)		
I-2-L	1	WD (0.27) (15.0-17.0')	2.2 (17.0-19.0')		
I-7-K	3	2.1 (8.5-10.5')	2.8 (8.5-10.5')		

- (1) Results of initial 201 samples; collected using top and bottom compositing technique.
 (2) Results of samples taken from the centers of archived Shelby tubes.
 (3) Results not reported elsewhere. Initial 201 sample taken by top and bottom compositing. Archive sample is from the archived Shelby tube center that initial sample was taken from.
 (4) Results not reported elsewhere.

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TABLE 5.6.1-1
SUMMARY OF MONITORING WELL DATA

WELL NUMBER	DEPTH TO STATIC WATER LEVEL ⁽¹⁾ (ft)	GROUND SURFACE ELEVATION ⁽²⁾ (ft)	DEPTH TO TOP OF SCREEN ⁽³⁾ (ft)	DEPTH TO BOTTOM OF SCREEN ⁽³⁾ (ft)	ESTIMATED SATURATED THICKNESS OPPOSITE WELL SCREEN (ft)	DEPTH INTERVAL (ft)
MW-1	6.6	98.7	3.5	14.2	7.9	6.6 to 14.5
MW-2	4.6	98.9	3.5	15.2	5.0	6.5 to 11.5
MW-3	4.7	97.3	3.0	8.5	3.8	4.7 to 8.5
MW-4	0.7	97.6	2.0	7.0	6.0	0.7 to 6.7
MW-5	4.2	98.9	3.0	8.5	3.8	4.2 to 8.0
MW-6	4.1	98.9	1.9	7.9	3.7	4.1 to 7.8
MW-7	1.6	98.4	2.0	8.2	1.9	1.6 to 3.5
MW-8	0.4	99.7	2.0	7.0	4.6	0.4 to 5.0

(1) Depths to static water level from ground surface at the time of slug tests (obtained from field log notebook).

(2) Elevations are with respect to site datum.

(3) Depths are with respect to ground surface.

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TABLE 5.6.1-2
RESULTS OF SLUG-TEST ANALYSIS

HYDRAULIC CONDUCTIVITY OF PERMEABLE
ZONE (ft/day)

WELL NUMBER	FALLING HEAD OR RISING HEAD	METHOD OF COOPER BREDEHOEFT & PAPADOPULOS	METHOD OF BOUWER AND RICE	
			Using Diam. Of Casing & Screen	Using Diam. of Drilled Hole
MW-1A	FH	8.3	9.1	74.3
	RH	17.6	7.5	60.9
MW-2A	RH	280	87.3	763
	RH	316	97.6	852
MW-3A	FH	42.6	56.0	404
	RH	14.6	21.2	152
MW-4A	FH	1.2	4.5	2.3
	RH	2.0	3.4	1.8
MW-5A	FH	11.6	20.4	152
	RH	48.5	22.8	168
MW-6A	RH	15.4	34.6	247
	FH	85.3	12.9	6.7
MW-7A	RH	62.2	15.8	126
	FH	7.0	11.7	6.0
MW-8A	RH	4.3	9.2	4.7

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TABLE 5.6.2-1
REPRESENTATIVE HYDRAULIC CONDUCTIVITY VALUES⁽¹⁾

WELL NUMBER	MEAN HYDRAULIC CONDUCTIVITY (ft/day)	RANGE IN HYDRAULIC CONDUCTIVITY (ft/day)	DEPTH RANGE OF MOST PERMEABLE ZONE (ft)
MW-1	10	5-15	6.6 to 14.5
MW-2	200	100-300	6.5 to 11.5
MW-3	40	20-60	4.7 to 8.5
MW-4	3	2-4	0.7 to 6.7
MW-5	20	10-30	4.2 to 8.0
MW-6	30	20-40	4.1 to 7.8
MW-7	10	5-15	1.6 to 3.5
MW-8	10	5-15	0.4 to 5.0

(1) Hydraulic conductivities are estimated values assigned to the zone in which the monitoring wells are screened. Results were determined from field slug tests.

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TABLE 5.6.4-1
GROUND WATER 2,3,7,8-TCDD REANALYSIS SUMMARY

SAMPLING DATE	INITIAL RESULT (ppb)	FINAL RESULT (ppb)
10-09-84	7.4	7.9
10-30-84	4.8 ⁽¹⁾	4.3
12-14-84	7.4	10.4

(1) A 100-ml sample from the original sampled water (no extract) after it
had been allowed to settle gave a 1.5 ppb result.

NOTE: All samples are from Well No. 2 and 5:1 dilution of the extract
was required.

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TABLE S.6.4-2
SUMMARY OF 2,3,7,8-TDDB
FOR GROUND WATER

WELL NUMBER	LOCATION	SAMPLING DATE	RESULTS (ppb)	SAMPLING DATE	RESULTS (ppb)	SAMPLING DATE	RESULTS (ppb)
1	I-2-L	10-09-84	0.68	10-30-84	0.56		
2	I-5-A	10-09-84	7.9	10-30-84	4.3	12-14-84	10.4
3	I-7-K	10-10-84	0.049	10-30-84	0.03		
4	C-7-C	10-09-84	0.20	10-30-84	0.74		
5	A-2-K	10-09-84	WD(0.008)	10-30-84	0.0059		
6	A-3-C	10-09-84	0.012	10-30-84	0.0086		
7	D-1-F	10-09-84	0.016	10-30-84	WD(0.024)		
8	F-7-B	10-09-84	0.72	10-30-84	1.1		

WD - not detected at the indicated () detection limit.

TABLE 5.6.4-3
SUMMARY OF DETECTED BASE/NEUTRAL/ACID ORGANICS
WELL WATER SAMPLES
(Expressed as µg/l or ppb)

	10-09-84			10-30-84		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
2,4,6-Trichlorophenol	1,700-11,000	3	8	290-3,900	3	8
2-Chlorophenol	290-4,600	3	8	11-3,600	4	8
2,4-Dichlorophenol	160-48,000	5	8	370-58,000	4	8
Phenol	36-3,700	5	8	43-600	3	8
Benzoic Acid	250	1	8	ND	0	8
2-Methylphenol	ND	0	8	24	1	8
4-Methylphenol	39-66	2	8	ND	0	8
2,4,5-Trichlorophenol	56-8,800	5	8	38-26,000	4	8
Acenaphthene	ND	0	8	30	1	8
1,2,4-Trichlorobenzene	200	1	8	9-890	3	8
Hexachlorobenzene	ND	0	8	770-860	2	8
2-Chloronaphthalene	ND	0	8	5	1	8
1,2-Dichlorobenzene	11-390	3	8	3-980	4	8
1,3-Dichlorobenzene	ND	0	8	13-200	2	8
1,4-Dichlorobenzene	110-590	3	8	6-1,200	4	8
Fluoranthene	15	1	8	3-120	5	8
Naphthalene	10-320	4	8	11-480	3	8
Bis(2-ethylhexyl)phthalate	55	1	8	3-75	3	8
Di-N-butylphthalate	12	1	8	8	1	8
Benzo(a)anthracene	ND	0	8	8	1	8
Anthracene	ND	0	8	4	1	8
Fluorene	10	1	8	32	1	8
Phenanthrene	2-34	2	8	3-110	5	8
Pyrene	3-19	3	8	5-46	5	8
Benzyl alcohol	8,000	1	8	4,300	1	8
2-Methylnaphthalene	7-260	4	8	3-900	6	8

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TABLE 5.6.4-4
SUMMARY OF DETECTED VOLATILE ORGANICS
WELL WATER SAMPLES
(Expressed as µg/l or ppb)

	10-09-84			10-30-84		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Benzene	3.0-3,900	8	8	10-7,900	7	8
Chlorobenzene	14-8,500	6	8	4-23,000	7	8
1,2-Dichloroethane	1,700	1	8	2,000	1	8
1,1,1-Trichloroethane	410	1	8	1,500	1	8
1,1-Dichloroethane	5	1	8	190	1	8
Chloroform	20-230	2	8	19-240	3	8
1,1-Dichloroethane	ND	0	8	53	1	8
trans-1,2-Dichloroethene	33-360	2	8	30-1,300	2	8
Ethylbenzene	44-740	3	8	43	2	8
Methylene chloride	6-12,000	8	8	3-7,400	8	8
Tetrachloroethene	2-5	2	8	2-43	3	8
Toluene	7-1,100	6	8	55-3,300	5	8
Trichloroethene	15-230	2	8	9-280	2	8
Vinyl chloride	28-88	2	8	24-220	2	8
Acetone	29-540	3	8	21-520	3	8
2-Butanone	870	1	8	180-430	2	8
Carbon disulfide	2-65	2	8	ND	0	8
4-Methyl-2-pentanone	3,300	1	8	1,800	1	8
Total xylenes	42-960	4	8	13-570	4	8

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TABLE 5.6.4-5
SUMMARY OF DETECTED HERBICIDES, PESTICIDES, AND PCB's
WELL WATER SAMPLES
(Expressed as ug/l or ppb)

	10-09-84			10-30-84		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
4,4'-DDT	17-22,000	4	8	14-2,770	4	8
4,4'-DDE	17-54	2	8	7-14	2	8
4,4'-DDD	15-13,000	5	8	7-1,390	4	8
Alpha-endosulfan	ND	0	8	1,240	1	8
2,4-D	6.9-27,000	6	8	74-20,000	4	8
2,4,5-T	470-5,600	4	8	68-3,500	4	8
2,4-DB	500	1	8	ND	0	8
Dinoseb (DNBP)	4.2	1	8	ND	0	8

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TABLE 5.6.4-6
SUMMARY OF DETECTED INORGANIC PARAMETERS
WELL WATER SAMPLES

(Expressed as ~~ug/l~~ or ~~ppb~~ *corrected per addenda (pg 5-2)*
ppm)

	10-09-84			10-30-84		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Antimony	0.003-0.151	7	8	0.001-0.024	8	8
Arsenic	0.015-0.621	8	8	0.028-0.629	8	8
Beryllium	0.003-0.008	5	8	0.002-0.010	7	8
Cadmium	0.002-0.029	8	8	0.002-0.023	8	8
Chromium	0.02-0.73	8	8	0.08-1.1	8	8
Copper	0.091-1.3	8	8	0.206-2.9	8	8
Lead	0.18-47	8	8	0.44-14	8	8
Mercury	0.001-0.16	8	8	0.002-0.066	8	8
Nickel	0.06-0.30	8	8	0.06-0.42	8	8
Selenium	ND	0	8	0.007	1	8
Silver	0.003-0.007	4	8	0.002-0.015	5	8
Zinc	0.247-17	8	8	0.864-17	8	8
Total Cyanide	0.01-0.35	7	8	0.01-0.63	7	8
Total Phenol	0.03-102	8	8	0.03-78	8	8

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TABLE 5.7.2-1
SUMMARY OF DETECTED COMPOUNDS IN
PASSAIC RIVER WATER SAMPLES
(Expressed as µg/l or ppb)

10/09/84			10/30/84		
CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
<u>VOLATILE ORGANICS</u>					
Chlorobenzene	ND	0	1	7	1
Chloroform	1	1	1	ND	1
trans-1,2-dichlorethane	2	1	1	2	1
Methylene chloride	4	1	1	7	1
Tetrachloroethene	3	1	1	3	1
Trichloroethene	2	1	1	1	1
Carbon disulfide	4	1	1	ND	1
<u>BASE/NEUTRAL/ACID ORGANICS</u>					
1,2-Dichlorobenzene	11	1	1	ND	1
Bis(2-ethylhexyl)phthalate	ND	0	1	2	1
Di-N-butylphthalate	6	1	1	ND	1
<u>HERBICIDES, PESTICIDES, AND PCB's</u>					
4,4'-DDT	3.5	1	1	ND	1
4,4'-DD	1.2	1	1	ND	1
<u>INORGANIC PARAMETERS</u>					
Arsenic	0.008	1	1	0.002	1
Chromium	ND	0	1	0.02	1
Copper	0.018	1	1	0.038	1
Nickel	0.10	1	1	ND	1
Zinc	0.011	1	1	0.049	1
Total Cyanide	0.02	1	1	0.01	1
Total Phenol	0.03	1	1	0.05	1

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TABLE 5.8-1
DESIGNATED ANALYSES FOR PASSAIC RIVER SEDIMENTS

STATION NUMBER	DEPTH	
	0-12 (inches)	12-24 (inches)
0-1-0	PP/D	NC
0-2-0	D	D
0-3-0	PP/D	NC
0-4-0	D	D
0-5-0	PP/D	PP/D
0-6-0	D	D
0-6-1	D	D
0-6-2	D	D
0-7-0	D	NC
0-8-0	PP/D	PP/D
0-8-1	PP/D	PP/D
0-8-2	PP/D	PP/D
0-9-0	PP/D	NC
1-0-0	PP/D	NC
1-1-0	D	D
1-1-1	D	D
1-1-2	D	D
1-2-0	D	D
1-3-0	PP/D	PP/D
1-4-0	D	NC
1-5-0	PP/D	NC
1-6-0	D	NC
1-7-0	D	NC

PP - priority pollutant defined by EPA contract laboratory program as acid/base/neutral, pesticide and PCB, and volatile organic compounds; herbicides, metals, total cyanides, total phenols.

D - Dioxin (2,3,7-8-TCDD).

NC - no sample designated to be collected.

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TABLE 5.8-2

TABLE 5.8-3
RESAMPLING 2,3,7,8-TCDD
RESULTS OF PASSAIC RIVER SEDIMENT STATION 1-3-0

SAMPLE IDENTIFICATION NUMBER	ELEVATION CODE	SAMPLE DEPTH	2,3,7,8-TCDD RESULTS (ppb)
1-3-0-1785-300	300	0-3'4"	151
1-3-0-1786-299	299	3'4"-3'10"	151
1-3-0-1787-298	298	3'10"-4'4"	176
1-3-0-1788-297	297	5'-5'6"	238
1-3-0-1789-296	296	5'6"-6'0"	450 ⁽¹⁾

(1) Reanalysis result from 1 gram sample aliquot; original (10 g) result
of 324 ppb was outside the linear calibration range.

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TABLE 5.8-4
PASSAIC RIVER SEDIMENT SAMPLES
ORGANIC PRIORITY POLLUTANT ANALYSIS LEVELS

STATION NUMBER	DEPTH (inches)	VOA	BASE/NEUTRAL/ACID	PESTICIDE	HERBICIDE
0-1-0	0-12	Low	Low	Low	Low
0-3-0	0-12	Low	Low	Low	Low
0-5-0	0-12	Low	Medium/Low ⁽¹⁾	Low	Low
0-5-0	12-24	Low	Medium/Low ⁽¹⁾	Low	Low
0-8-0	0-12	Low	Medium/Low ⁽¹⁾	Low	Low
0-8-0	12-24	Low	Low	Low	Low ⁽²⁾
0-8-1	0-12	Low	Low	Low	Low
0-8-1	12-24	Low	Low	Low	Low
0-8-2	0-12	Low	Medium	Low	Low
0-8-2	12-24	Low	Low	Low	Low ⁽²⁾
0-9-0	0-12	Low	Low	Low	Low
1-0-0	0-12	Low	Low ⁽³⁾	Low	Low
1-3-0	0-12	Low	Low	Medium	Low
1-3-0	12-24	Low	Medium	Medium	Low ⁽⁴⁾
1-5-0	0-12	Low	Low	Low	Low ⁽²⁾

(1) These samples contained only one B/N/A compound at a concentration greater than the medium level detection limit; they were re-extracted and analyzed as low-level samples.

(2) Further diluted 1:5 - See Organics Analysis Data Sheet.

(3) Further diluted 1:2 - See Organics Analysis Data Sheet.

(4) Further diluted 1:200 - See Organics Analysis Data Sheet.

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0-12 INCHES				12-24 INCHES			
CONCENTRATION RANGE	RESULTS	NUMBER	ANALYZED	CONCENTRATION RANGE	RESULTS	NUMBER	ANALYZED
2,4-Dichlorophenol	-	0	10	360,000	1	1	5
2,4,5-Trichlorophenol	-	0	10	140,000	1	1	5
Acenaphthene	24,000	1	10	1,200	1	1	5
Fluoranthene	78,000-300	8	10	1,800-610	4	4	5
Naphthalene	-	0	10	710	1	1	5
Bis(2-ethylhexyl)phthalate	66,000-12,000	9	10	37,000-11,000	4	4	5
Di-N-octylphthalate	640-230	2	10	860-590	2	2	5
Benzo(a)anthracene	42,000-810	2	10	590	1	1	5
Benzo(a)pyrene	30,000	1	10	880	1	1	5
Benzo(b)fluoranthene	29,000	1	10	780	1	1	5
Chrysene	95,000	1	10	3,400-1,200	2	2	5
Acenaphthylene	8,300-440	2	10	360	1	1	5
Anthracene	58,000	1	10	1,700-330	2	2	5
Fluorene	18,000	1	10	1,500	1	1	5
Phenanthrene	110,000-440	4	10	3,900-430	3	3	5
Pyrene	100,000-260	6	10	3,600-520	4	4	5
2-Methylnaphthalene	250	1	10	1,600	1	1	5

TABLE 5.8-5
SUMMARY OF DETECTED BASE/NEUTRAL/ACID ORGANIC COMPOUNDS
PASSAIC RIVER SEDIMENTS
(Expressed as ug/kg or ppb)

TABLE 5.8-6
SUMMARY OF DETECTED VOLATILE ORGANIC COMPOUNDS
PASSAIC RIVER SEDIMENTS
(Expressed as $\mu\text{g/kg}$ or ppb)

	0-12 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Benzene	28-7	6	10	210	1	5
Chlorobenzene	250-53	4	10	96	1	5
Methylene chloride	640-73	10	10	680-65	5	5
Chloromethane	-	0	10	140	1	5
Tetrachloroethane	-	0	10	22	1	5
Toluene	32	1	10	270-52	2	5
Acetone	1,600-220	8	10	830-190	5	5
2-Butanone	160-70	4	10	310-69	4	5
Carbon disulfide	31-9	3	10	25-12	2	5
Total xylenes	400-140	2	10	500	1	5

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TABLE 5.8-7
SUMMARY OF DETECTED HERBICIDE, PESTICIDES AND PCBs
PASSAIC RIVER SEDIMENTS
 (Expressed as µg/kg or ppb)

	0-12 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Dieldrin	-	0	10	12,000	1	5
4-4'-DDT	4,400-430	4	10	1,300	1	5
4,4'-DDE	93-23	5	10	75-24	3	5
4,4'-DDD	58-22	6	10	350-44	2	5
PCB-1242	720-120	8	10	8,100-460	3	5
PCB-1254	300-200	2	10	200	1	5
Dalapon	180-120	2	10	-	0	5
Dichloroprop (2,4-DP)	470	1	10	-	0	5
2,4-D	900-130	5	10	490,000	1	5
2,4,5-T	100-81	3	10	820,000-76	2	5
Dinoseb (DNBP)	300-180	6	10	-	0	5

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TABLE 5.8-8
SUMMARY OF DETECTED INORGANIC PARAMETERS
PASSAIC RIVER SEDIMENTS
(Expressed as ug/kg or ppb)

	0-12 INCHES			12-24 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
Antimony	3.3-0.3	10	10	3.4-0.2	5	5
Arsenic	79-7.5	10	10	97-8.3	5	5
Beryllium	1.1-0.39	10	10	0.85-0.54	5	5
Cadmium	21.0-4.8	10	10	16-3	5	5
Chromium	970-200	10	10	550-260	5	5
Copper	700-220	10	10	720-320	5	5
Lead	760-410	10	10	700-460	5	5
Mercury	18.0-4.9	10	10	13-3	5	5
Nickel	116-51	10	10	114-55	5	5
Silver	11.0-4.0	10	10	9.4-4.2	5	5
Zinc	2,100-700	10	10	1,500-850	5	5
Total Cyanide	4.5-<0.5	10	10	6.8-1.02	5	5
Total Phenols	1.5-0.02	8	10	298-0.4	5	5

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TABLE 5.9.1-1
RESULTS OF 2,3,7,8-TCDD ANALYSIS
OF BORING SAMPLES FROM SHERWIN-WILLIAMS PROPERTY
BROWN STREET AND LISTER AVENUE
NEWARK, NEW JERSEY

ELEVATION CODE	DEPTH (inches)	RESULTS (ppb)
100	0-6	1.2
101	6-12	5.1
102	12-24	3.4
109	above silt (11-12.5')	ND(0.57)
201	silt (15-17')	ND(0.76)

ND - not detected at the indicated () detection limit.

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0497

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TABLE 5.9.2-1
RESULTS OF 2,3,7,8-TCDD ANALYSIS
NEWARK, NEW JERSEY BACKGROUND SAMPLES
(Expressed as ug/kg or ppb)

LOCATION	DEPTH (inches)	RESULTS
Harrison Avenue	0-6	ND(0.17)
Raymond Boulevard	0-6	ND(0.27)
Roanoke Avenue	0-6	ND(0.77)

ND - not detected at the indicated () detection limit.

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TABLE 5.9.2-2
SUMMARY OF DETECTED BASE/NEUTRAL/ACID ORGANIC COMPOUNDS
NEAR-SURFACE SOILS
FROM SITE AND MEANBY BACKGROUND SAMPLES
(Expressed as µg/kg or ppb)

	SITE - 0-6 INCHES			SITE - 12-24 INCHES			MEANBY BACKGROUND - 0-6 INCHES		
	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	NUMBER SAMPLES ANALYZED
2,4,6-Trichlorophenol	3,500,000-1,300	5	21	1,700,000-8,700	4	21	-	0	3
2,4-Dichlorophenol	3,600,000-900	7	21	2,500,000-870	0	21	-	0	3
2,4-Dimethylphenol	-	0	21	1,700,000	1	21	-	0	3
Benzoic Acid	1,800	1	21	-	0	21	-	0	3
2,4,5-Trichlorophenol	15,000,000-870	5	21	7,500,000-2,500	5	21	-	0	3
Acenaphthene	250	1	21	-	0	21	-	0	3
1,2,4-Trichlorobenzene	17,000-1,500	2	21	19,000	1	21	-	0	3
Hexachlorobenzene	110,000-560	13	21	720,000-3,200	6	21	620,000-110,000	2	3
1,2-Dichlorobenzene	520-230	2	21	9,000	1	21	-	0	3
1,3-Dichlorobenzene	-	0	21	610	1	21	-	0	3
1,4-Dichlorobenzene	1,400-470	3	21	1,300	1	21	-	0	3
Fluorant hene	6,100-330	5	21	64,000-670	6	21	3,500-2,400	3	3
Heptachlorene	700	1	21	0,200	1	21	400	1	3
Bis(2-ethylhexyl)phthalate	1,300-110	3	21	310,000-3,100	3	21	1,700-670	3	3

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0500

REMARK BACKGROUND - 0-6 INCHES				80 LISTER - 12-24 INCHES				80 LISTER - 0-6 INCHES			
NUMBER	POSITIVE	CONCENTRATION	ANALYZED	NUMBER	POSITIVE	CONCENTRATION	ANALYZED	NUMBER	POSITIVE	CONCENTRATION	ANALYZED
1	1	200	21	2	370,000-2,000	21	0	-	0	21	220
3	3	1,900-1,500	21	5	47,000-510	21	3	47,000-910	3	21	47,000-910
3	3	1,500-1,200	21	5	44,000-560	21	3	4,800-1,000	3	21	4,800-1,000
3	3	2,700-2,200	21	5	71,000-960	21	3	7,100-2,100	3	21	7,100-2,100
3	3	3,700-3,200	21	6	120,000-1,400	21	2	12,000-2,600	2	21	12,000-2,600
3	3	610-250	21	2	860-240	21	2	690-210	2	21	690-210
3	3	600-500	21	3	1,200-620	21	4	3,000-310	4	21	3,000-310
3	3	2,300-1,500	21	1	32,000	21	3	11,000-3,300	3	21	11,000-3,300
3	3	2,800-1,300	21	2	300-230	21	1	320	1	21	320
3	3	-	21	6	61,000-440	21	5	4,100-230	5	21	4,100-230
3	3	1,700-1,100	21	2	21,000-400	21	2	2,500-2,200	2	21	2,500-2,200
3	3	1,700-1,400	21	7	70,000-200	21	6	2,200-230	6	21	2,200-230
3	3	-	21	1	430	21	0	-	0	21	Pyrene
3	3	-	21	1	21,000	21	1	21,000	1	21	2-Methylanthracene
3	3	-	21	1	21,000	21	1	21,000	1	21	2-Methylanthracene

TABLE 5.9.2-2
(Continued)

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0501

SITE - 0-6 INCHES				SITE - 12-24 INCHES				NEWMARK BACKGROUND - 0-6 INCHES			
CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER
Benzene	21	1	21	23,000-11	3	21	21	-	0	3	3
Chlorobenzene	84,000-19	2	21	170,000-22	6	21	21	-	0	3	3
Chloroform	18	1	21	38,000-13	2	21	21	-	0	3	3
Ethylbenzene	-	0	21	60,000	1	21	21	-	0	3	3
Methylbenzene chloride	1,500-14	21	21	130,000-21	21	21	21	66-32	3	3	3
Tetrachloroethane	860	1	21	36,000-1,100	2	21	21	-	0	3	3
Toluene	-	0	21	2,000,000-7	6	21	21	-	0	3	3
Trichloroethane	-	0	21	9	1	21	21	-	0	3	3
Acetone	5,000-56	13	21	2,000-68	15	21	21	-	0	3	3
2-Butanone	1,400-130	2	21	9,200-51	6	21	21	-	0	3	3
Carbon disulfide	-	0	21	1	1	21	21	-	0	3	3
2-Heptanone	-	0	21	36,000	1	21	21	-	0	3	3
Total aliphatics	-	0	21	310,000	1	21	21	-	0	3	3

TABLE 3.9.2-3
SUMMARY OF DETECTED VOLATILE ORGANICS
NEAR-SURFACE SOILS
FROM SITE AND NEWMARK BACKGROUND SAMPLES
(Expressed as mg/kg or ppb)

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0502

SITE - 0-6 INCHES				SITE - 12-24 INCHES				REMARK BACKGROUND - 0-6 INCHES			
CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER	CONCENTRATION RANGE	NUMBER POSITIVE RESULTS	ANALYZED SAMPLES	NUMBER
1,500,000-620	19	21	15	5,090,000-1,400	15	21	21	200	1	3	3
93,000-20	9	21	8	37,000-1,200	8	21	21	77-32	2	3	3
13,000-1,700	3	21	5	166,000-1,200	5	21	21	-	0	3	3
8,900	1	21	1	1,400	1	21	21	-	0	3	3
-	0	21	0	-	0	21	21	1,700-1,200	2	3	3
70,000-190	9	21	9	29,000-420	9	21	21	-	0	3	3
7,600-740	10	21	13	85,000-190	13	21	21	-	0	3	3
2,100-190	9	21	10	80,000-490	10	21	21	-	0	3	3

4,4'-DOT
4,4'-DDE
4,4'-DDD
Alpha-Endosulfan
PCB-1260
Dieldrin

TABLE 3.9.2-4
SUMMARY OF DETECTED HERBICIDES, PESTICIDES AND PCBs
FROM SITE AND REMARK BACKGROUND SAMPLES
(Expressed as ug/kg or ppb)

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0503

SITE - 0-6 INCHES				SITE - 12-24 INCHES				NEWARK BACKGROUND - 0-6 INCHES			
ANALYZED	RESULTS	POSITIVE	CONCENTRATION RANGE	ANALYZED	RESULTS	POSITIVE	CONCENTRATION RANGE	ANALYZED	RESULTS	POSITIVE	CONCENTRATION RANGE
NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS	NUMBERS
Antimony	14	21	0.0-0.09	17	21	0.0-0.10	21	21	3	3	9.1-2.2
Argon	21	21	23-0.13	21	21	41-0.60	21	21	3	3	10-4.6
Beryllium	11	21	0.03-0.22	9	21	0.04-0.25	21	21	2	3	0.5-0.47
Cadmium	12	21	2.0-0.09	14	21	2.0-0.08	21	21	3	3	2.0-2.0
Chromium	21	21	50-1.1	21	21	50-3.9	21	21	3	3	90-51
Copper	21	21	200-2.4	20	21	250-2.0	21	21	3	3	311-127
Lead	21	21	607-1.0	21	21	646-2.1	21	21	3	3	1,700-595
Mercury	10	21	39-0.1	10	21	37-0.4	21	21	3	3	2.0-0.6
Nickel	20	21	02-3.1	20	21	40-2.1	21	21	3	3	74-35
Selenium	1	21	0.48	3	21	2.2-0.01	21	21	0	3	-
Silver	7	21	1.2-0.24	6	21	11-0.23	21	21	3	3	1.4-0.45
Zinc	21	21	20,000-20	21	21	1,300-0.0	21	21	3	3	020-420
Total Cyanide	19	21	1.97-0.15	10	21	2.0-0.10	21	21	3	3	2.0-0.70
Total Phenols	20	21	0.7-0.0.20	21	21	3,370-0.10	21	21	1	3	117

TABLE 3.9.2-5
SUMMARY OF DETECTED INORGANIC PARAMETERS
FROM SITE AND NEWARK BACKGROUND SAMPLES
(Expressed as ug/kg or ppb)

TABLE 5.10-1
2,3,7,8-TCDD ANALYSIS RESULTS
DRUM SAMPLING PROGRAM

DIOXIN CONCENTRATION (ppb)	SAMPLE ID	SAMPLE DESCRIPTION	NUMBER OF DRUMS REPRESENTED BY THIS SAMPLE
12.1	0018-0045-D-L	Drum No. 18, CY	15
12,220	0021-0064-D-L	white and yellow crystals Drum No. 21, CQ	11
8.0	0040-0091-D-L	yellow crystall powders Drum No. 40, 23AA	5
54.0	0065-0136-D-L	milky liquid Drum No. 65, 400,	37
2.6	0075-0152-D-L	clear gold liquid Drum No. 75, 15T	9
13.9	0119-0255-D-L	pink thick liquid Drum No. 119, CZ	13
1.5	0162-0346-D-L	dark brown liquid Drum No. 162, CX	8
35.9	0176-0364-D-L	golden liquid Drum No. 176, 21Y	89
16.0	0183-0371-D-L	thick white paste Drum No. 183, QQ	14
7.5	0174-0403-D-L	pink and red liquid Drum No. 174, 21Y	89
3.4	0230-0502-D-L	thick white paste Drum No. 230, BB	31
476	0251-0523-D-L	clear liquid and white solids Drum No. 251, ZB	32
ND (1.7)	0305-0670-D-L	brown sludge and water Drum No. 305, Pit	11
ND (6.7)	0314-0679-D-L	clear liquid Drum No. 314, 9K	6
ND (3.8)	0388-0816-D-L	dark brown crystals Drum No. 388, 18W	43
ND (2.0)	0392-0820-D-L	clear liquid (rusty) Drum No. 392, JJ	38
12	0438-0925-D-L	golden liquid Drum No. 438, MM	13
ND (16.2)	0450-0937-D-L	white solids Drum No. 450, DD	12
174	0458-0948-D-L	white powder Drum No. 458, S	4
ND (8.4)	0492-1015-D-L	brown liquid Drum No. 492, PP	17
ND (2.0)	0554-1136-D-L	dark liquids w/solids Drum No. 554, Pit 3	89
8,750	0558-1140-D-L	clear liquid Drum No. 558, Pit 3	89
		dark sludge w/water	

ND = none detected; number in () indicates the lower detection limit of the linear range due to background noise.

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TABLE 5.11-1
RESULTS OF ANALYSES OF SELECTED SAMPLES
2,3,7,8-TCDD, 2,3,7,8-TCDF, AND OCDD

SAMPLE IDENTIFICATION	TYPE OF SAMPLE	2,3,7,8-(1) TCDD (ppb)	2,3,7,8-(2) TCDF (ppb)	OCDD(3)
H-2-H-1598-100	NSSS	93.5	1.9	81
A-3-C-0354-101	BSS	18.8	5.0	22.0
I-2-L-0857-109	BSS	12.1	12.6	82.0
B-2-M-1346-102	NSSS	2.8	ND(0.36)	ND(0.88)
A-5-C-1661-101	NSSS	453	ND(0.66)	4.8
F-5-E-1605-101	NSSS	394	10.6	10.0
C-5-E-1567-101	NSSS	217	217	37
H-7-H-1521-101	NSSS	27.6	25	3.3
H-5-F-1395-101	NSSS	69.3	ND(0.60)	8.7
A-2-C-1334-101	NSSS	330	2.3	38
C-7-C-0642-100	NSSS	180	8.2	49
A-2-K-0436-102	BSS	72.5	0.70	70
C-7-C-0710-109	BSS	71.8	1.0	2.2
I-7-K-1047-109	BSS	5.8	10.1	ND(0.62)
I-5-A-0869-109	BSS	20.9	ND(0.96)	27
C-7-C-0701-201	BSS	2.1	ND(0.49)	1.1
9600-1834-108	BSS-	3.4	19.9	15.7
Sherwin-Williams				
C-5-F-1448-100	NSSS	361	13.2	24
I-3-O-0309-299	River Sediment	130	14.9	16
I-1-O-0302-299	River Sediment	65.6	0.67	5.6
I-3-O-1785-300	River Sediment- Bed Sampling	151	11.4	10.6
O-9-O-0299-300	River Sediment	10.8	ND(0.43)	4.8
O-1-O-0186-300	River Sediment	3.9	ND(0.69)	8.5
O-8-O-0205-299	River Sediment	10.4	ND(0.23)	8.6

(1) 2,3,7,8-Tetrachlorodibenzo-p-dioxin.

(2) 2,3,7,8-Tetrachlorodibenzofuran.

(3) Octachlorodibenzo-p-dioxin.

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TABLE 5.12-1
QUALITY ASSURANCE OBJECTIVES

ANALYSIS	MATRIX	METHOD PRECISION (% Rel. Std. Dev.)	METHOD ACCURACY (% Recovery)
2,3,7,8-TCDD	Water	±25	50-120
	Soil	±25	50-120
	Air/Ind.Hyg.	±25	50-120
	Surface Wipes	±25	50-120
	Misc. Organics	±25	50-120
Volatile Priority Pollutants	Water	±7-22 ⁽¹⁾	75-120 ⁽¹⁾
	Soil	±7-22 ⁽¹⁾	60-130 ⁽¹⁾
	Ambient Air	±7-22 ⁽¹⁾	60-130 ⁽¹⁾
	Misc. Organics	±7-22 ⁽¹⁾	60-130 ⁽¹⁾
Semi-volatile Priority Pollutants	Water	±6-70 ⁽¹⁾	40-180 ⁽¹⁾
	Soil	±6-70 ⁽¹⁾	40-180 ⁽¹⁾
	Ambient Air	±6-70 ⁽¹⁾	40-180 ⁽¹⁾
	Misc. Organics	±6-70 ⁽¹⁾	40-180 ⁽¹⁾
Priority Pollutant Metals	Water	±10 ⁽¹⁾	70-100 ⁽¹⁾
	Soil	±15 ⁽¹⁾	70-100 ⁽¹⁾
	Misc. Organics		
Cyanides	Water	±10	85
	Soil	±15	80
Total Phenols	Water	±1	75-100
	soil	±5	75-100

(1) The accuracy and precision are compound dependent.

NOTES: The Quality Assurance Objective for completeness is 90 percent for all of the above analyses.

The Method Precision and Method Accuracy reported for each analysis were generated by the EPA under ideal conditions. The precision and accuracy that can be achieved are frequently determined by the level of interferences present rather than instrumental or method limitations.

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TABLE 5.12.1-1
FIELD BLANK COLLECTION SUMMARY

MATRIX	NUMBER SAMPLES COLLECTED	ANALYSIS PARAMETERS (SAMPLES)	NUMBER BLANKS COLLECTED	ANALYSIS PARAMETERS (BLANKS)	PERCENT FREQUENCY (BLANKS TO SAMPLES)
Solid	148	Full PP* (87)	35	VOA	24
		or	2	Dioxin	1.4
		Dioxin only (61)	2	Extractable PP	1.4
			2	PP Metals	1.4
			2	Cyanide	1.4
			2	Phenols	1.4
Water	19	Full PP*	2	Full PP*	10.5
Wipes	78	Dioxin only	15	Dioxin	15
Chips	70	Dioxin only	2	Dioxin	2.9

*PP indicates priority pollutants which includes dioxin.

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TABLE 5.12.1-2
QUALITY CONTROL RESULTS
FIELD BLANKS FOR 2,3,7,8-TCDD ANALYSIS
(Concentration Units are in ppb)

SAMPLE TYPE/NUMBER	COLLECTION DATE	RESULT
Field Blank/L1556	10-15-84	ND(0.0007)
Field Blank/L1623	10-17-84	ND(0.009)

ND = not detected, detection limit reported in parentheses.

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TABLE 5.12.1-3
QUALITY CONTROL RESULTS
FIELD AND TRIP BLANKS FOR PRIORITY POLLUTANT METALS ANALYSIS
(Concentration Units are mg/L - ppm)

PARAMETER	FIELD BLANK (K1374) 10-09-84*	TRIP BLANK (K1375) 10-09-84*	FIELD BLANK (K1703) 10-19-84*
Antimony	<0.001**	<0.001	<0.001
Arsenic	<0.001	<0.001	0.004
Beryllium	<0.001	<0.001	<0.002
Cadmium	<0.001	<0.001	<0.001
Chromium	<0.01	<0.01	<0.01
Copper	<0.002	<0.002	0.005
Lead	<0.01	<0.01	<0.01
Mercury	<0.001	0.001	<0.001
Nickel	<0.01	<0.01	<0.01
Selenium	<0.001	<0.001	<0.001
Silver	<0.002	<0.002	<0.002
Thallium	<0.02	<0.02	<0.02
Zinc	<0.001	<0.001	0.012

*Dates shown are collection dates.

**The less-than result indicates a value below the presented detection limit.

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TABLE 5.12.1-4
QUALITY CONTROL RESULTS
FIELD AND TRIP BLANKS FOR CYANIDE/PHENOLS ANALYSIS
(Concentration Units are in Mg/L - ppm)

SAMPLE TYPE/NUMBER	COLLECTION DATE	PARAMETER	RESULT
Field Blank/K1429	10-10-84	Cyanide	<0.01*
Trip Blank/K1430	10-10-84	Cyanide	<0.01
Field Blank/K1659	10-18-84	Cyanide	<0.01
Field Blank/K1513	10-11-84	Phenols	<0.01
Trip Blank/K1514	10-11-84	Phenols	<0.01
Field Blank/K1732	10-22-84	Phenols	0.08

*The less-than result indicates a value below the presented detection limit.

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TABLE 5.12.1-5
SUMMARY OF VOLATILE CONTAMINANTS
IN FIELD AND TRIP BLANKS

COMPOUND	FIELD BLANKS		TRIP BLANKS	
	NUMBER SAMPLES DETECTED	CONCENTRATION RANGE (ppb)	NUMBER SAMPLES DETECTED	CONCENTRATION RANGE (ppb)
Methylene chloride	37	(5-130)	37	(5-51)
1,1,1-Trichloroethane	1	4	0	-
Chloroform	19	(1-61)	6	(1-54)
Tetrachloroethene	1	4	0	-
Toluene	2	(1-2)	2	(1-1)
Trichloroethene	1	5	0	-
Acetone	5	(10-47)	7	(10-78)
Bromodichloromethane	14	(3-12)	4	(9-14)
Carbon disulfide	1	2	0	-

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TABLE 5.12.2.1-1
QUALITY CONTROL ACCEPTANCE CRITERIA
SURROGATE RECOVERIES

SURROGATE COMPOUND	ACCEPTABLE PERCENT RECOVERY RANGE*	
	WATER	SOIL
VOA:		
1,2-Dichloroethane-d4	77-120	64-129
Toluene-d8	86-119	69-127
4-Bromofluorobenzene	85-121	61-122
B/M/A:		
Nitrobenzene-d5	41-120	24-115
2-Fluorobiphenyl	44-119	37-120
p-Terphenyl-d14	33-128	28-133
2-Fluorophenol	23-107	24-111
Phenol-d5	15-96	20-106
2,4,6-Tribromophenol	20-105	11-102
Pesticide:		
Dibutyl chlorendate	67-114	0-205

*In accordance with EPA contract laboratory program requirements.

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TABLE 5.12.2.1-2
QUALITY CONTROL ACCEPTANCE CRITERIA
SPIKE RECOVERIES AND RPD

SURROGATE COMPOUND	ACCEPTABLE PERCENT RECOVERY RANGE		ACCEPTABLE [*] RPD (%)
	WATER	SOIL	WATER/SOIL
VOA:			
1,1-Dichloroethylene	61-145	59-177	<15
Trichloroethylene	71-120	62-137	<15
Benzene	76-127	66-142	<15
Toluene	76-125	59-139	<15
Chlorobenzene	75-130	60-133	<15
B/W/A:			
1,2,4-Trichlorobenzene	39-98	38-107	<50
Acenaphthene	46-118	31-137	<50
2,6-Dinitrotoluene	24-96	28-89	<50
Di-n-butylphthalate	11-117	29-135	<50
Pyrene	26-127	35-142	<50
N-Nitrosodi-n-propylamine	41-116	41-126	<50
1,4-Dichlorobenzene	36-97	28-104	<50
Pentachlorophenol	9-103	17-109	<40
4-Chloro-3-methylphenol	23-97	26-103	<40
Phenol	12-89	26-90	<40
2-Chlorophenol	27-123	25-102	<40
4-Nitrophenol	10-80	11-114	<40
Pesticide			
Lindane	56-123	46-127	<40
Heptachlor	40-131	35-130	<40
Aldrin	40-120	34-132	<40
Dieldrin	52-126	31-134	<40
Endrin	56-121	42-139	<40
p,p'-DDT	38-127	23-134	<40

*In accordance with EPA contract laboratory program requirements.

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ANALYSIS	SURROGATE COMPOUND	AVERAGE PERCENT RECOVERY (Std.Dev.)			
		LOW	MEDIUM	BASE/NEUTRAL/ACID	PESTICIDE
Toluene-d8	99 (26.4)	94 (27.3)	-	-	-
4-Bromofluorobenzene	95 (26.4)	97 (26.8)	-	-	-
1,2-Dichloroethane-d4	103 (23.6)	104 (26.0)	-	-	-
Nitrobenzene-d5	-	-	75 (218)	78 (222)	-
2-Fluorobiphenyl	-	-	76 (213)	83 (212)	-
p-Terphenyl-d14	-	-	68 (227)	62 (223)	-
Phenol-d5	-	-	75 (215)	83 (212)	-
2-Fluorophenol	-	-	71 (216)	77 (211)	-
2,4,6-Tribromophenol	-	-	70 (222)	79 (220)	-
Isobutyl chloroacetate	-	-	-	-	100 (17.5) 104 (27.9)

TABLE 5.12.2.1.1-1
QC SUMMARY DATA
ORGANIC PRIORITY POLLUTANT SURROGATE
RECOVERY RESULTS
SOILS

TABLE 5.12.2.1.1-2
 QUALITY CONTROL SUMMARY DATA:
 ORGANIC PRIORITY POLLUTANT SURROGATE RECOVERY RESULTS
 OUTSIDE OF QUALITY CONTROL ACCEPTANCE LIMITS

ANALYSIS SURROGATE COMPOUND	NUMBER OUTLIERS/TOTAL NUMBER ANALYSES					
	LOW VOLATILE	MEDIUM VOLATILE	LOW BASE/NEUTRAL/ACID	MEDIUM BASE/NEUTRAL/ACID	LOW PESTICIDE	MEDIUM PESTICIDE
Toluene-d8	0/68	0/7	-	-	-	-
4-Bromofluorobenzene	0/68	0/7	-	-	-	-
1,2-Dichloroethane-d4	3/68	0/7	-	-	-	-
Nitrobenzene-d5	-	-	2/46	0/29	-	-
2-Fluorobiphenyl	-	-	0/46	0/29	-	-
p-Terphenyl-d14	-	-	3/46	0/29	-	-
Phenol-d5	-	-	2/46	0/29	-	-
2-Fluorophenol	-	-	1/46	0/29	-	-
2,4,6-Tribromophenol	-	-	2/46	1/29	-	-
Dibutyl chlorodate	-	-	-	-	32/43	19/32

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TABLE 5.12.2.1.1-3
QUALITY CONTROL CHECK FREQUENCIES
SOILS

ANALYSIS/LEVEL	NUMBER SAMPLES ANALYZED	NUMBER MS/MSD PAIRS ANALYZED	QC CHECK FREQUENCY (%)
VOA/Low	68	5	7.4
VOA/Medium	7	2	29
BNA/Low	46	4	8.7
BNA/Medium	29	2	6.9
Pesticide/Low	43	4	9.3
Pesticide/Medium	32	2	6.3
Herbicide	75	5	7.5

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TABLE 5.12.2.1.1-4
 QUALITY CONTROL SUMMARY DATA
 VOLATILE ORGANIC PRIORITY POLLUTANT SPIKE RECOVERY
 AND DUPLICATE RESULTS
 SOILS

SPIKE COMPOUND	LOW VOA ANALYSIS		MEDIUM VOA ANALYSIS	
	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)
1,1-Dichloroethylene	101 (\pm 9.4)	4.8 (\pm 3.9)	101 (\pm 15)	5.5 (\pm 1.5)
Trichloroethylene	86 (\pm 6.8)	4.6 (\pm 5.7)	90 (\pm 3.5)	6.0 (\pm 3.0)
Benzene	87 (\pm 6.7)	4.0 (\pm 4.0)	79 (\pm 7.5)	4.5 (\pm 0.5)
Toluene	90 (\pm 12)	7.4 (\pm 4.9)	106 (\pm 24.9)	8.5 (\pm 3.5)
Chlorobenzene	85 (\pm 4.3)	4.8 (\pm 6.0)	89 (\pm 4.2)	6.5 (\pm 6.5)

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SPIKE COMPOUND	LOW RNA ANALYSIS			MEDIUM RNA ANALYSIS		
	AVERAGE % RECOVERY (1 Std.Dev.)	AVERAGE RPD (1Std.Dev.)	AVERAGE % RECOVERY (1Std.Dev.)	AVERAGE RPD (1Std.Dev.)	AVERAGE % RECOVERY (1Std.Dev.)	AVERAGE RPD (1Std.Dev.)
1,2,4-Trichlorobenzene	70 (112)	3.3 (13.4)	99 (15.1)	6.0 (12.0)	9.5 (19.5)	2.0 (12.0)
Acenaphthene	77 (17.5)	11 (15.9)	88 (18.2)	9.5 (19.5)	2.0 (12.0)	4.0 (12.0)
2,6-Dinitrofluorene	52 (15.3)	6.3 (17.2)	59 (14.6)	2.0 (12.0)	4.0 (12.0)	8.0 (12.0)
Di-n-butylphthalate	83 (112)	11 (113)	91 (112)	4.0 (12.0)	8.0 (12.0)	132 (17.0)
Pyrene	77 (19.9)	6.8 (15.0)	132 (17.0)	8.0 (12.0)	6.0 (11.0)	3.0 (11.0)
N-nitrosodimethylamine	75 (111)	7.0 (19.0)	94 (113)	6.0 (11.0)	3.0 (11.0)	3.0 (11.0)
1,4-Dichlorobenzene	81 (15.3)	4.5 (15.0)	90 (11.6)	3.0 (11.0)	3.0 (11.0)	12 (10.5)
Pentachlorophenol	71 (115)	12 (19.1)	100 (12.6)	3.0 (11.0)	3.0 (11.0)	1.5 (11.5)
4-Chloro-3-methylphenol	51 (17.3)	5.8 (11.3)	73 (16.0)	12 (10.5)	4.0 (10)	90 (12.1)
Phenol	70 (16.1)	9.8 (11.8)	88 (10)	4.0 (10)	4.0 (10)	16 (10.5)
2-Chlorophenol	73 (16.8)	11 (11.9)	90 (12.1)	4.0 (10)	4.0 (10)	16 (10.5)
4-Nitrophenol	55 (125)	11 (19.9)	56 (139)	16 (10.5)	16 (10.5)	16 (10.5)

TABLE 5.12.2.1.1-5
QUALITY SUMMARY DATA
BASE/NEUTRAL/ACID PRIORITY POLLUTANT SPIKE
RECOVERY AND DUPLICATE RESULTS
SOILS

TABLE S.12.2.1.1-6
 QUALITY CONTROL SUMMARY DATA
 PRIORITY POLLUTANT, PESTICIDE, AND HERBICIDE SPIKE RECOVERY
 AND DUPLICATE RESULTS
 SOILS

SPIKE COMPOUND	LOW PESTICIDE ANALYSIS		MEDIUM PESTICIDE ANALYSIS		HERBICIDE ANALYSIS	
	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)
Lindane	99 (\pm 10)	6.0 (\pm 3.5)	103 (\pm 2.0)	1.5 (\pm 1.5)	-	-
Heptachlor	100 (\pm 13)	9.0 (\pm 3.1)	111 (\pm 9.0)	7.5 (\pm 7.5)	-	-
Aldrin	96 (\pm 8.2)	6.5 (\pm 3.2)	103 (\pm 12)	12 (\pm 12)	-	-
Dieldrin	100 (\pm 14)	7.0 (\pm 3.5)	110 (\pm 3.6)	3.5 (\pm 0.5)	-	-
Endrin	95 (\pm 5.9)	3.3 (\pm 2.4)	107 (\pm 3.9)	6.0 (\pm 3.0)	-	-
p,p'-DDT	107 (\pm 13)	18 (\pm 10)	NM	3.5 (\pm 3.5)	-	-
Alaapon	-	-	-	-	49 (\pm 39)	21 (\pm 12)
Dicamba	-	-	-	-	71 (\pm 36)	3.5 (\pm 2.0)
Dichloroprop	-	-	-	-	74 (\pm 38)	7.5 (\pm 2.6)
2,4-D	-	-	-	-	61 (\pm 76)	21 (\pm 17)
2,4,5-TP	-	-	-	-	65 (\pm 33)	4.4 (\pm 2.6)
2,4,5-T	-	-	-	-	45 (\pm 26)	16 (\pm 15)
2,4-DM	-	-	-	-	49 (\pm 29)	49 (\pm 38)
Dimethab	-	-	-	-	0	NM

NM = Not reportable = all percent recoveries outside quality control limits.
 NI = not calculatable = all recoveries are zero.

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MA - no sediments were analyzed as medium-level VOCs.

ANALYSIS	SURROGATE COMPOUND	LOW VOLATILE	MEDIUM VOLATILE	LOW BNA	MEDIUM BNA	LOW PESTICIDE	MEDIUM PESTICIDE	AVERAGE PERCENT RECOVERY (±STD.DEV.)
	Toluene-d8	102 (±2.2)	NA	-	-	-	-	
	4-Bromofluorobenzene	95 (±8.2)	NA	-	-	-	-	
	1,2-Dichloroethane-d4	101 (±2.0)	MA	-	-	-	-	
	Nitrobenzene-d5	-	-	92 (±4.7)	93 (±3.3)	-	-	
	2-Fluorobiphenyl	-	-	67 (±4.1)	71 (±8.5)	-	-	
	p-Terphenyl-d14	-	-	75 (±9.9)	71 (±21)	-	-	
	Phenol-d5	-	-	85 (±8.3)	88 (±9.3)	-	-	
	2-Fluorophenol	-	-	79 (±8.5)	80 (±8.4)	-	-	
	2,4,6-Tribromophenol	-	-	85 (±10)	83 (±12)	-	-	
	Dibutyl chlorododecyl	-	-	-	-	-	-	111 (±8.6) 111 (±9.0)

TABLE 5.12.2.1.2-1
QUALITY CONTROL SUMMARY DATA
ORGANIC PRIORITY POLLUTANT SURROGATE
RECOVERY RESULTS
SEDIMENTS

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TABLE 5.12.2.1.2-2
QUALITY CONTROL SUMMARY DATA
VOLATILE PRIORITY POLLUTANT AND HERBICIDE SPIKE RECOVERY
AND DUPLICATE RESULTS
SEDIMENTS

SPIKE COMPOUND	LOW VOLATILE ANALYSIS		LOW HERBICIDE ANALYSIS	
	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD*	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD*
1,1-Dichloroethylene	150 (\pm 10)	13	-	-
Trichloroethylene	95 (\pm 5)	11	-	-
Benzene	115 (\pm 5)	9	-	-
Toluene	115 (\pm 5)	9	-	-
Chlorobenzene	115 (\pm 5)	9	-	-
Dalapon	-	-	62 (\pm 5)	6
Dicamba	-	-	82 (\pm 2)	2
Dichloroprop	-	-	80 (\pm 5.5)	8
2,4-D	-	-	95 (\pm 12)	12
2,4,5-TP	-	-	73 (\pm 3)	0
2,4,5-T	-	-	69 (\pm 4.5)	19
2,4,5-DB	-	-	47 (\pm 29)	118
Dinoseb	-	-	2.1 (\pm 1.8)	48

*Only one matrix spike/matrix spike duplicate pair was run for each analysis type; therefore, no average RPD is available.

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*Only one MS/MSD pair was run for each analyte type; therefore, no average RPD is available.
NR = not reportable - both recoveries are outside quality control limits.

SPIKE COMPOUND	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD (%)	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD (%)
1,2,4-Trichlorobenzene	95 (25)	10	93 (22)	4
Acenaphthene	78 (21)	2	79 (24)	10
2,6-Dinitrotoluene	60 (21)	3	53 (22)	7
Di-n-butylphthalate	88 (20)	0	87 (21)	2
Pyrene	NR	0	75 (20)	0
M-nitrosodipropylamine	95 (21)	2	100 (27)	14
1,4-Dichlorobenzene	77 (20)	0	80 (21)	2
Pentachlorophenol	76 (22.5)	6	85 (21)	2
4-Chloro-3-methylphenol	74 (22.5)	7	80 (21.5)	4
Phenol	83 (20)	0	84 (23)	7
2-Chlorophenol	83 (20)	0	79 (24)	10
4-Mitrophenol	23 (20.5)	7	25 (20.5)	4

TABLE 5.12.2.1.2-3
QUALITY CONTROL SUMMARY DATA
BASE/NEUTRAL/ACID PRIORITY POLLUTANT SPIKE RECOVERY
AND DUPLICATE RESULTS
SEDIMENTS

TABLE 5.12.2.1.2-4
 QUALITY CONTROL SUMMARY DATA
 PRIORITY POLLUTANT PESTICIDE SPIKE RECOVERY
 AND DUPLICATE RESULTS
 SEDIMENTS

SPIKE COMPOUND	LOW PESTICIDE ANALYSIS		MEDIUM PESTICIDE ANALYSIS	
	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD ^a (%)	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD ^a (%)
Lindane	90 (± 2.5)	7	102 (± 4)	8
Heptachlor	105 (± 1.5)	3	106 (± 3)	6
Aldrin	95 (± 2)	4	100 (± 2.5)	6
Dieldrin	94 (± 2)	4	78 (± 1.5)	7
Endrin	112 (± 2)	3	120 (± 1.5)	3
p,p'-DDT	107 (± 3.5)	7	117 (± 1)	2

^aOnly one MS/MSD pair was run for each analysis type; therefore, no average RPD is available.

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TABLE 5.12.2.1.3-1
QUALITY CONTROL SUMMARY DATA
ORGANIC PRIORITY POLLUTANT SURROGATE
RECOVERY RESULTS
WATERS

ANALYSIS SURROGATE COMPOUND	AVERAGE % RECOVERY (\pm STD.DEV.)		
	LOW VOLATILE	LOW BNA	LOW PESTICIDE
Toluene-d8	102 (\pm 4.2)	-	-
4-Bromofluorobenzene	99 (\pm 4.6)	-	-
1,2-Dichloroethane-d4	103 (\pm 4.3)	-	-
Nitrobenzene-d5	-	80 (\pm 12)	-
2-Fluorobiphenyl	-	62 (\pm 9.3)	-
p-Terphenyl-d14	-	46 (\pm 10)	-
Phenol-d5	-	62 (\pm 14)	-
2-Fluorophenol	-	50 (\pm 11)	-
2,4,6-Tribromophenol	-	55 (\pm 15)	-
Dibutyl chlorendate	-	-	89 (\pm 18)

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TABLE 5.12.2.1.3-2
QUALITY CONTROL SUMMARY DATA
ORGANIC PRIORITY POLLUTANT SURROGATE
WATER RECOVERY RESULTS
OUTSIDE OF QUALITY CONTROL
ACCEPTANCE LIMITS

ANALYSIS SURROGATE COMPOUND	NUMBER OF OUTSIDE RECOVERIES/TOTAL NUMBER OF ANALYSES		
	LOW VOLATILE	LOW BNA	LOW PESTICIDE
Toluene-d8	0/24	-	-
4-Bromofluorobenzene	0/24	-	-
1,2-Dichloroethane-d4	0/24	-	-
Nitrobenzene-d5	-	8/24	-
2-Fluorobiphenyl	-	8/24	-
p-Terphenyl-d14	-	8/24	-
Phenol-d5	-	7/24	-
2-Fluorophenol	-	7/24	-
2,4,6-Tribromophenol	-	8/24	-
Dibutyl chloredate	-	-	18/24

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TABLE 5.12.2.1.3-3
QUALITY CONTROL DATA
ORGANIC PRIORITY POLLUTANT SPIKE RECOVERY
AND DUPLICATE RESULTS
WATERS

SPIKE COMPOUND	LOW VOLATILE ANALYSIS		LOW BNA ANALYSIS		LOW PESTICIDE ANALYSIS	
	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)	AVERAGE % RECOVERY (\pm Std.Dev.)	AVERAGE RPD (\pm Std.Dev.)	AVERAGE % RECOVERY (\pm Std.Dev.)	RPD* (%)
1,1-Dichloroethylene	110 (\pm 4.1)	2 (\pm 0)	-	-	-	-
Trichloroethylene	103 (\pm 8.4)	5 (\pm 1)	-	-	-	-
Benzene	98 (\pm 2.6)	5 (\pm 1)	-	-	-	-
Toluene	104 (\pm 6.3)	10 (\pm 3.5)	-	-	-	-
Chlorobenzene	102 (\pm 5.0)	9 (\pm 4.5)	-	-	-	-
1,2,4-Trichlorobenzene	-	-	63 (\pm 4.1)	3.5 (\pm 3.5)	-	-
Acenaphthene	-	-	62 (\pm 1.9)	5.5 (\pm 2.5)	-	-
2,6-Dinitrotoluene	-	-	47 (\pm 14)	6.5 (\pm 0.5)	-	-
Di-n-butylphthalate	-	-	37 (\pm 2.7)	13 (\pm 3.5)	-	-
Styrene	-	-	59 (\pm 16)	15 (\pm 6.5)	-	-
N-nitrosodi-n-propylamine	-	-	70 (\pm 11)	3.5 (\pm 3.5)	-	-
1,4-Dichlorobenzene	-	-	55 (\pm 1.2)	2 (\pm 2)	-	-
2,4-Dichlorophenol	-	-	73 (\pm 3.9)	7.5 (\pm 0.5)	-	-
2-Chloro-3-methylphenol	-	-	54 (\pm 7.9)	4 (\pm 1)	-	-
Phenol	-	-	66 (\pm 11)	6 (\pm 2)	-	-
2-Chlorophenol	-	-	72 (\pm 9.2)	7 (\pm 0)	-	-
2-Nitrophenol	-	-	65 (\pm 7.5)	6.5 (\pm 1.5)	-	-
Lindane	-	-	-	-	72 (\pm 1.5)	4
Heptachlor	-	-	-	-	68 (\pm 5)	14
Aldrin	-	-	-	-	69 (\pm 4)	11
Dieldrin	-	-	-	-	75 (\pm 2.5)	7
Endrin	-	-	-	-	81 (\pm 1.5)	4
p,p'-DDE	-	-	-	-	78 (\pm 3.5)	9

*Only one MS/MSD pair was run for this analysis type; therefore, an average RPD is not available.

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TABLE 5.12.2.2-1
QUALITY CONTROL CHECK SAMPLE FREQUENCIES
INORGANIC/CLASSICAL ANALYSES

SAMPLE MATRIX	TOTAL NUMBER OF SAMPLES ANALYZED	NUMBER OF SAMPLE SPIKES	PERCENT FREQUENCY (%)	NUMBER OF BLIND SPLITS	PERCENT FREQUENCY (%)
Soil	75	5	6.7	5	6.7
Sediment	15	1	6.7	1	6.7
Water	24	2	8.3	2	8.3

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TABLE 5.12.2.2-2
QUALITY CONTROL SUMMARY DATA
INORGANIC PARAMETERS SPIKE RECOVERY AND
DUPLICATE RESULTS

MATRIX PARAMETER	AVERAGE \bar{x} RPD (\pm STD.DEV.)		AVERAGE \bar{x} RPD (\pm STD.DEV.)
	WATER	SOIL	WATER AND SOIL
Antimony	133 (\pm 33)	24 (\pm 27)	105 (\pm 31)
Arsenic	70 (\pm 33)	28 (\pm 20)	113 (\pm 25)
Beryllium	27 (\pm 2)	9.8 (\pm 10)	102 (\pm 7.3)
Cadmium	20 (\pm 13)	15 (\pm 14)	95 (\pm 7.1)
Chromium	74 (\pm 74)	15 (\pm 10)	94 (\pm 7.0)
Copper	16 (\pm 11)	20 (\pm 16)	102 (\pm 7.0)
Lead	16 (\pm 8)	20 (\pm 17)	105 (\pm 6.8)
Mercury	60 (\pm 60)	43 (\pm 37)	107 (\pm 24)
Nickel	11 (\pm 4)	28 (\pm 16)	112 (\pm 6.5)
Selenium	0	0	92 (\pm 19)
Silver	9 (\pm 9)	3.7 (\pm 8.2)	99 (\pm 6.7)
Thallium	0	0	104 (\pm 7.8)
Zinc	13 (\pm 10)	13 (\pm 5)	105 (\pm 7.9)

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TABLE 5.12.2.3-1
LABORATORY DUPLICATE RESULTS SUMMARY
2,3,7,8-TCDD

SAMPLE TYPE	ORIGINAL RESULT (ppb)	DUPLICATE RESULT (ppb)	RPD (%)
Chip	2.0	1.8	10
Chip	ND (0.08)*	ND(0.23)	0
Sediment	0.53	0.55	3.7
Chip	9.2	14	40
Sediment	0.60	1.1	58
Chip	ND(0.28)	ND(0.54)	0
Sediment	1.7	1.4	19
Soil	ND(0.30)	ND(0.09)	0
Bulk	3.0	2.5	18
Chip	76.8	77.5	0.9
Chip	43.2	58.5	30
Chip	18.1	18.1	0
Chip	0.93	1.0	7.3
Soil	2.5	1.7	38
Soil	2.4	1.2	70
Soil	5.8	4.6	23
Soil	2.2	2.9	27
Soil	193.	214	10
Soil	0.39	0.30	26
Soil	217.	223	2.7
Soil	7.3	6.1	18
Soil	3.4	3.3	3.0
Soil	268.	257.	4.2
Soil	33.4	26	25
Sediment	450.	646	36
Soil	0.27	0.16	51
Drum	10,200.	9150.	11

*The numbers in parentheses represent the detection limits.

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TABLE 5.12.2.3-2
LABORATORY SPIKE RECOVERY RESULTS SUMMARY
DIOXIN

SAMPLE TYPE	ORIGINAL RESULT (ppb)	+	AMOUNT SPIKED (ppb)	=	THEORET. CONC SAMPLE + SPIKE (ppb)	SPIKE RESULT (ppb)	PERCENT RECOVERY
Chip	ND		1.0		1.0	0.77	77
Sediment	0.53		1.0		1.5	1.3	87
Sediment	0.60		1.0		1.6	1.4	88
Chip	ND		1.0		1.0	1.0	100
Sediment	1.7		1.0		2.7	2.2	82
Soil	ND		1.0		1.0	1.2	120
Chip	0.93		1.0		1.9	2.4	126
Soil	2.2		1.0		3.2	2.7	84
Soil	0.39		1.0		1.4	1.6	114
Soil	0.27		17.0		17.3	18.1	105
Soil	2.5		1.0		3.5	2.5	71
Soil	2.4		1.0		3.4	2.4	71
Soil	5.8		1.0		6.8	6.5	96
Soil	3.4		1.0		4.4	3.5	80

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TABLE 5.12.3.1-1
PROGRAM QC BLANK SAMPLE RESULTS
2,3,7,8-TCDD ANALYSIS

SAMPLE NUMBER	MATRIX	DATE SHIPPED TO LAB	RESULT
9900-1458-W-L	Wipe	10-15-84	ND (3.6 ng/wipe)
9900-1459-W-L	Wipe	10-15-84	(2)
9900-1460-W-L	Wipe	10-15-84	ND(3.8 ng/wipe)
Q-1-C-1464-100-S-L	Soil	10-15-84	0.76 ppb
Q-1-C-1465-100-S-L	Soil	10-15-84	1.6 ppb
Q-1-C-1466-100-S-L	Soil	10-15-84	0.89 ppb
9900-1470-W-Y	Water	10-19-84	ND(0.002 ppb)

(1) Detection limit is cited in the parentheses.

(2) Sample lost during preparation - no results available.

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TABLE 5.12.3.2-1
PROGRAM QC BLANK SPIKE SAMPLE RESULTS
2,3,7,8-TCDD ANALYSIS

SAMPLE NUMBER	MATRIX	DATE ASSIGNED	SPIKE LEVEL	RESULT	% RECOVERY
9900-1461-W-L	Wipe	10-15-84	40 ng/wipe	34.9 ng/wipe	87.
9900-1462-W-L	Wipe	10-15-84	40 ng/wipe	34.7 ng/wipe	87.
9900-1463-W-L	Wipe	10-15-84	40 ng/wipe	38.5 ng/wipe	96.
Q-1-C-1467-100-S-L	Soil	10-15-84	671. ppb	725. ppb	108.
Q-1-C-1468-100-S-L	Soil	10-15-84	671. ppb	878. ppb	131.
Q-1-C-1469-100-S-L	Soil	10-15-84	671. ppb	780. ppb	116.

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TABLE 5.12.3.3-1
ITAS SAMPLE SPLIT RESULTS SUMMARY
2,3,7,8-TCDD ANALYSIS

ORIGINAL/SPLIT SAMPLE NOS.	MATRIX	ORIGINAL RESULT (ppb)	SPLIT RESULT (ppb)	RPD (ppb)
L0111/L0112	Chip	0.70	0.95	30
L0890/L0891 ⁽¹⁾	Chip	1.25	1.20	4
L0894/L0895 ⁽¹⁾	Chip	ND (0.78) ⁽⁴⁾	ND (0.48)	0
L1039/L1049	Soil	3510. ⁽²⁾	824. ⁽²⁾	120
		1500. ⁽³⁾	1300. ⁽³⁾	14
L1517/L1519	Soil	1.2	0.7	53
L1661/L1663	Soil	453. ⁽²⁾	526. ⁽²⁾	15
		460. ⁽³⁾	440. ⁽³⁾	4
Y1038/Y1048	Soil	350. ⁽²⁾	1030. ⁽²⁾	98
		350. ⁽³⁾	449. ⁽³⁾	25
Y1333/Y1339	Soil	326. ⁽²⁾	462. ⁽²⁾	34
		296. ⁽³⁾	249. ⁽³⁾	18
Y1471/Y1474	Soil	3.6	1.8	66
Y1359/Y1371	Water	0.68	0.18	116

(1) These samples were collected from the Sergeant site, concurrently with the 80 Lister Avenue sampling.

(2) One gram sample aliquot result.

(3) Ten gram sample aliquot result.

(4) Detection limit indicated in parentheses.

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TABLE 5.12.4-1
NJDEP-DESIGNATED SPLIT SAMPLES

ITC SAMPLE NUMBER	SAMPLE TYPE	ITAS REQUIRED ANALYSES	NJDEP SPLIT SAMPLE NUMBER
1-1-0-0301-300-M-L	Sediment	Dioxin	9300-0286-300-M-J
1-6-9-0312-300-M-L	Sediment	Dioxin	9300-0287-300-M-J
4506-0425-C-L	Chip	Dioxin	9300-0429-C-J
4506-0426-C-L	Chip	Dioxin	9300-0430-C-J
0265-0573-D-L	Drum	HazCat	ETC No. F5470A
0266-0574-D-L	Drum	HazCat	ETC No. F5471A
0267-0575-D-L	Drum	HazCat	ETC No. F5472A
0268-0576-D-L	Drum	HazCat	ETC No. F5473A
0269-0577-D-L	Drum	HazCat	ETC No. F5474A
0270-0578-D-L	Drum	HazCat	ETC No. F5475A
0271-0579-D-L	Drum	HazCat	ETC No. F5476A
0272-0580-D-L	Drum	HazCat	ETC No. F5477A
0273-0581-D-L	Drum	HazCat	ETC No. F5478A
0274-0582-D-L	Drum	HazCat	ETC No. F5479A
0275-0583-D-L	Drum	HazCat	ETC No. F5480A
0276-0584-D-L	Drum	HazCat	ETC No. F5481A
0277-0585-D-L	Drum	HazCat	ETC No. F5482A
0278-0586-D-L	Drum	HazCat	ETC No. F5483A
0279-0587-D-L	Drum	HazCat	ETC No. F5484A
0280-0588-D-L	Drum	HazCat	ETC No. F5485A
0281-0589-D-L	Drum	HazCat	ETC No. F5486A
0282-0590-D-L	Drum	HazCat	ETC No. F5487A
0437-0924-D-L	Drum	HazCat	ETC No. F5488A
0438-0925-D-L	Drum	HazCat/Dioxin	ETC No. F5489A
0439-0926-D-L	Drum	HazCat	ETC No. F5490A
0440-0927-D-L	Drum	HazCat	ETC No. F5491A
0441-0928-D-L	Drum	HazCat	ETC No. F5492A
0442-0929-D-L	Drum	HazCat	ETC No. F5493A
0443-0930-D-L	Drum	HazCat	ETC No. F5494A
0444-0931-D-L	Drum	HazCat	ETC No. F5495A
0450-0937-D-L	Drum	HazCat/Dioxin	No ETC No. recorded
0451-0938-D-L	Drum	HazCat	No ETC No. recorded
0452-0939-D-L	Drum	HazCat	No ETC No. recorded
0453-0940-D-L	Drum	HazCat	No ETC No. recorded
1-7K-1038-100-S-Y	Soil	All parameters	ETC No. 6506 & 6508
8012-1323-Z-L	Sewer	Dioxin	ETC No. F5467A
C-5-F-1448-100-S-Y	Soil	All parameters	ETC No. F7608A & F7606E
A-4-F-1516-100-S-Y	Soil	All parameters	ETC No. F7603E
A-4-F-1517-101-S-L	Soil	Dioxin	No ETC No. recorded
C-3-I-1577-101-S-L	Soil	Dioxin	No ETC No. recorded

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TABLE 5.12.4-2
NJDEP SOIL PROFICIENCY SAMPLE RESULTS

CLIENT NUMBER	SORT	SAMPLE DESCRIPTION	RESULTS
9400-1549-S-L	841012	A010-Blank Spike	1.7 ppb
9400-1550-S-L	841012	A011	1.2 ppb
9400-1551-S-L	841012	A012	3.6 ppb
9400-1552-S-L	941012	A013	492 ppb
9400-1592-S-L	841015	A016	502 ppb
9400-1593-S-L	841015	A017	541 ppb
9400-1594-S-L	841015	A018	1.1 ppb
9400-1595-S-L	841015	A019-Blank Spike	1.5 ppb
9400-1475-S-L	841016	A020-Blank Spike	1.7 ppb
9400-1476-S-L	841016	A021	4.4 ppb
9400-1477-S-L	841016	A022	1.1 ppb
9400-1478-S-L	841016	A023	511 ppb
9400-1653-S-L	841017	A024-Blank Spike	1.7 ppb
9400-1654-S-L	841017	A025	4.2 ppb
9400-1655-S-L	841017	A026	1.2 ppb
9400-1656-S-L	841017	A027	393 ppb
9400-1675-S-L	841018	A028-Blank Spike	1.8 ppb
9400-1676-S-L	841018	A029	4.5 ppb
9400-1677-S-L	841018	A030	1.4 ppb
9400-1678-S-L	841018	A031	595 ppb
9400-1733-S-L	841019	A032-Blank Spike	1.6 ppb
9400-1734-S-L	841019	A033	MD (0.76 ppb)
9400-1735-S-L	841019	A034	554 ppb
9400-1736-S-L	841029	A035	500 ppb
9400-1762-S-L	841022	A036-Blank Spike	1.9 ppb
9400-1781-S-L	841023	A038-Blank Spike	1.6 ppb
9400-1782-S-L	841023	A039	367 ppb

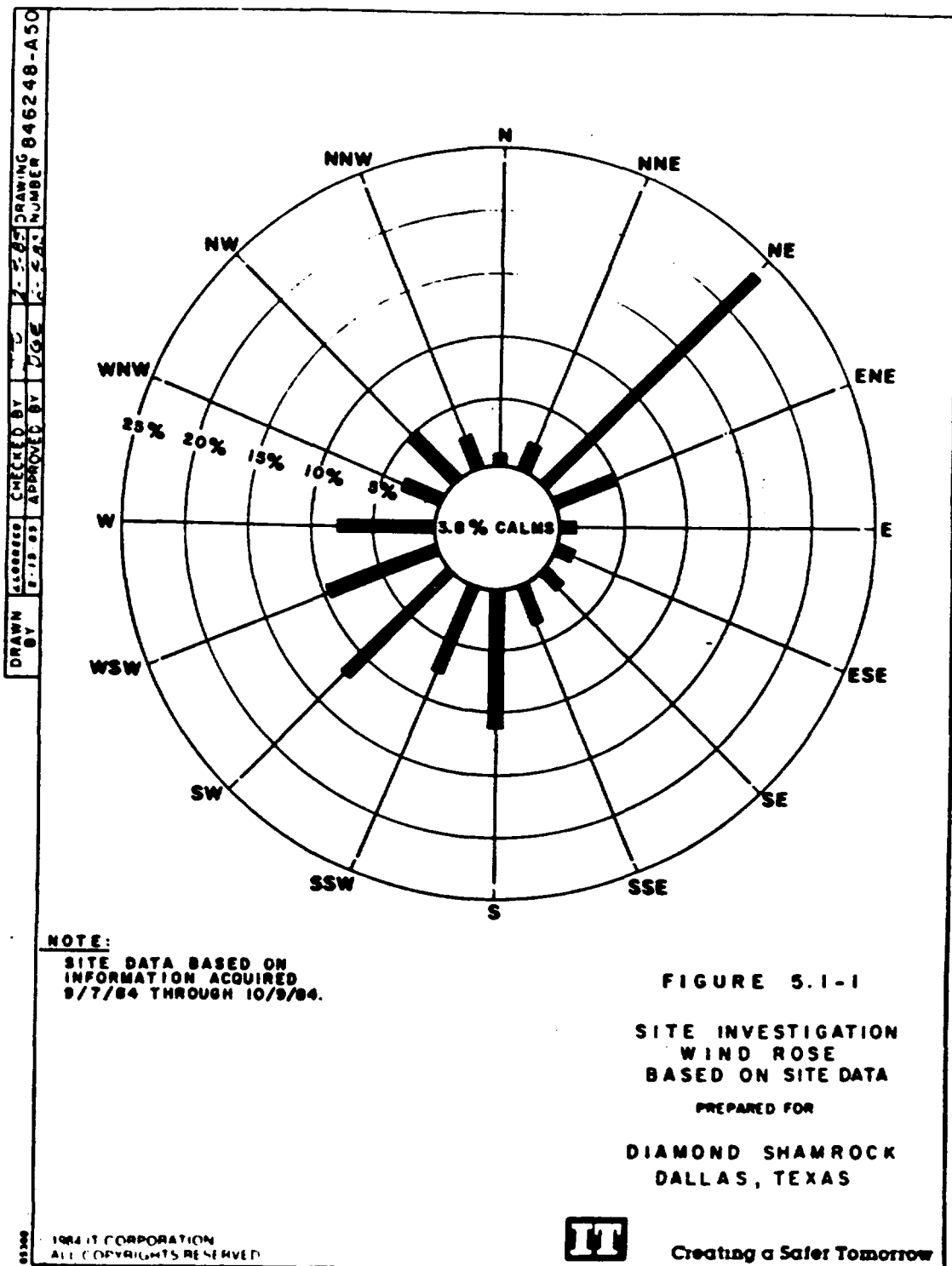
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FIGURES

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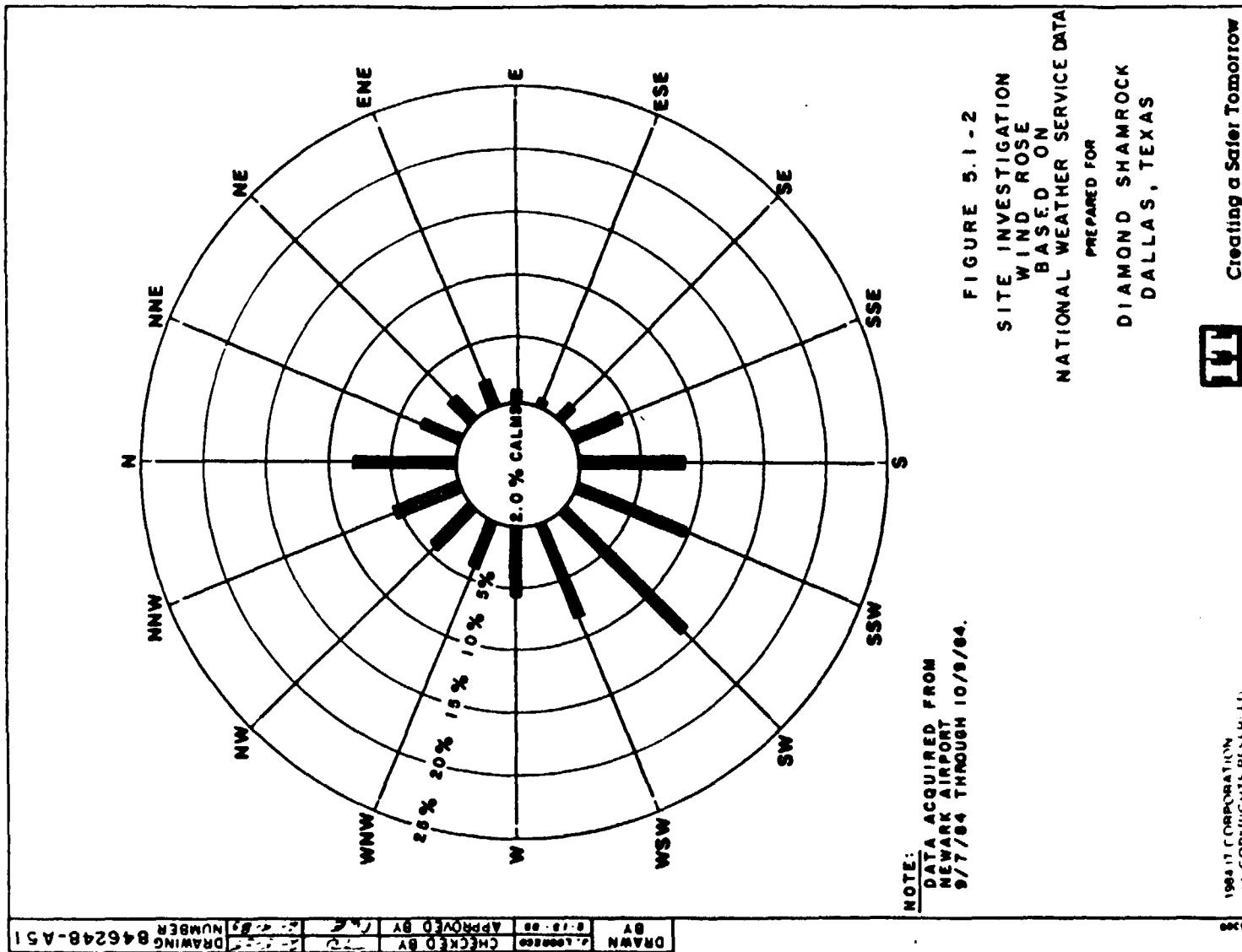
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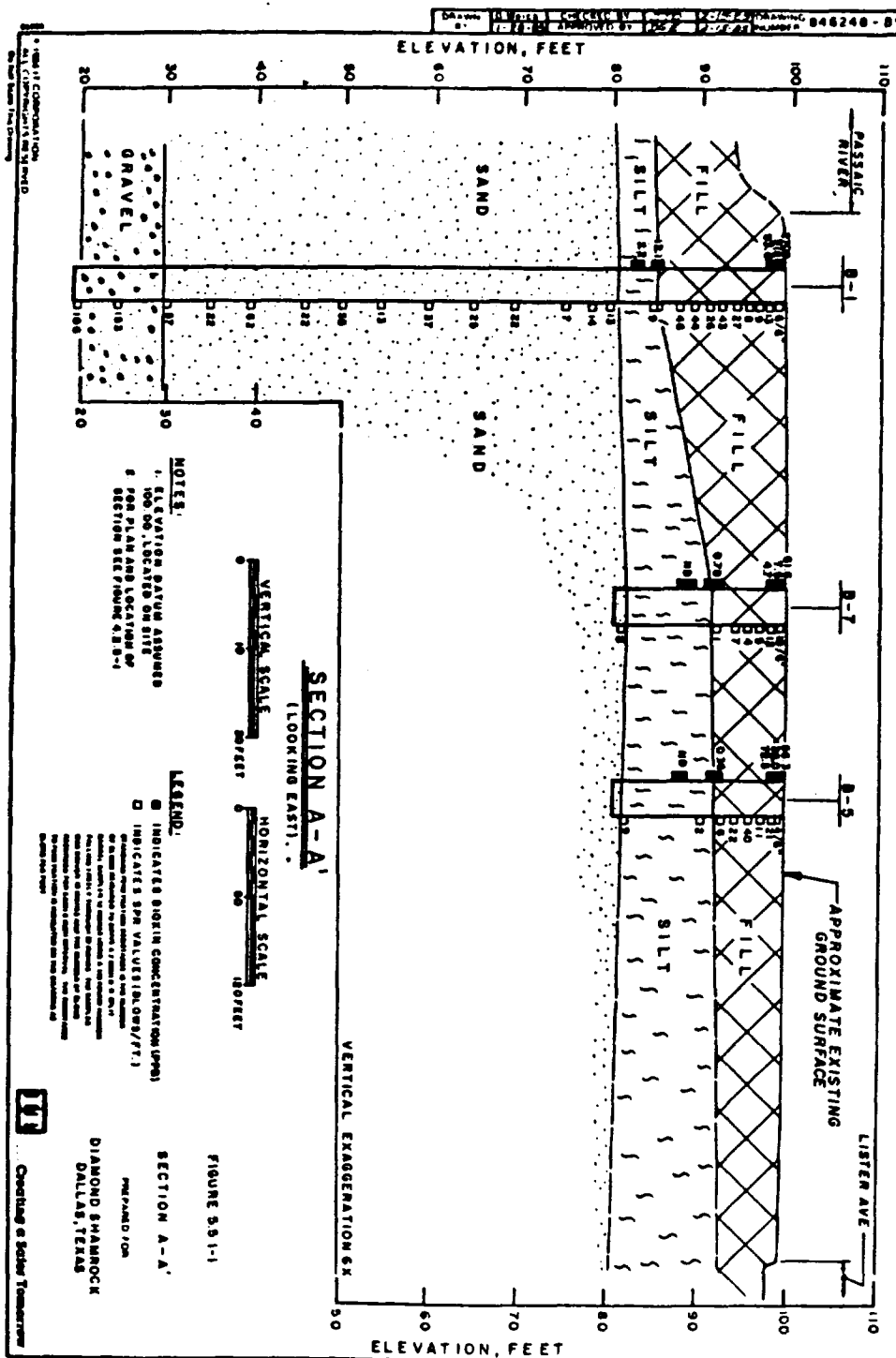
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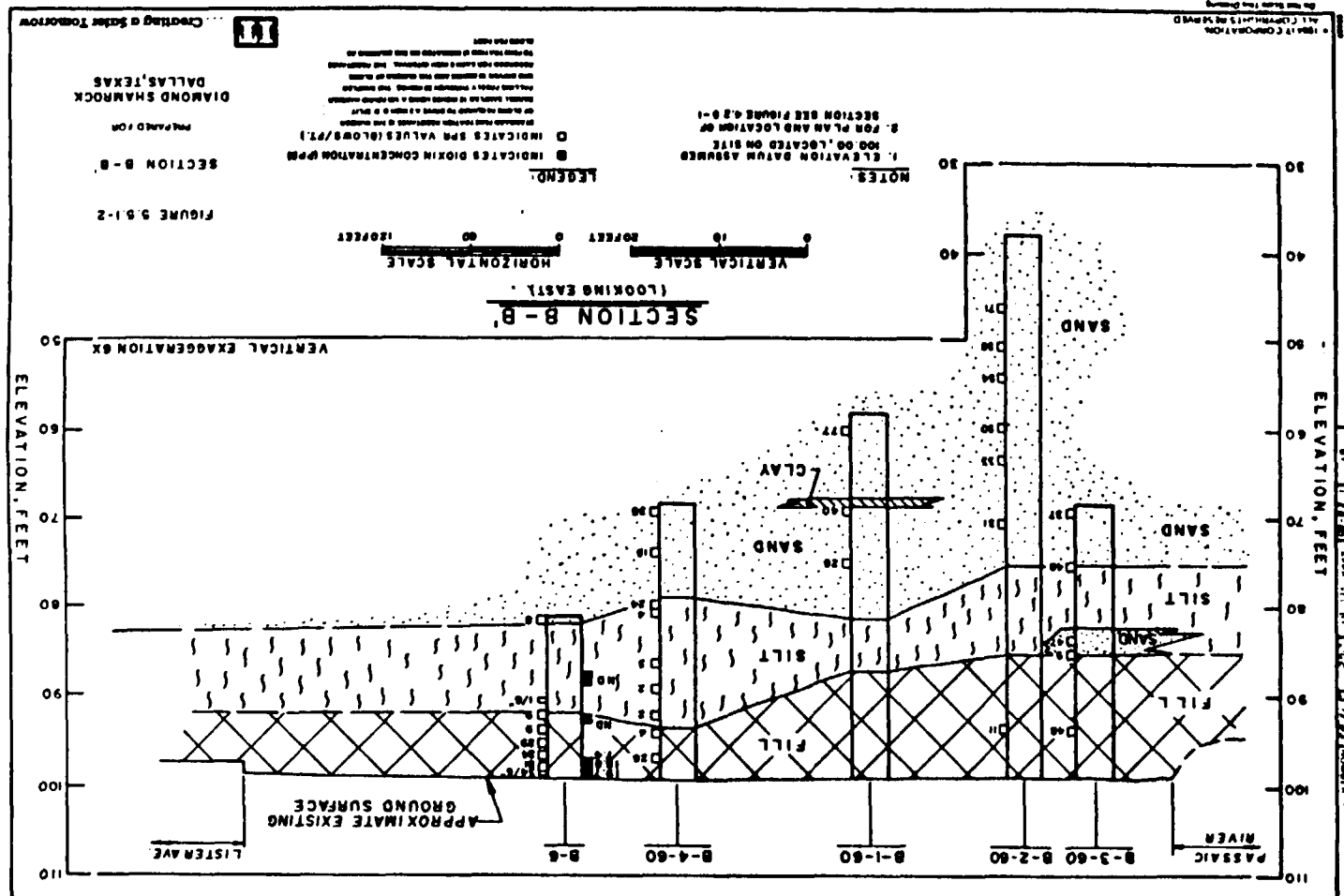


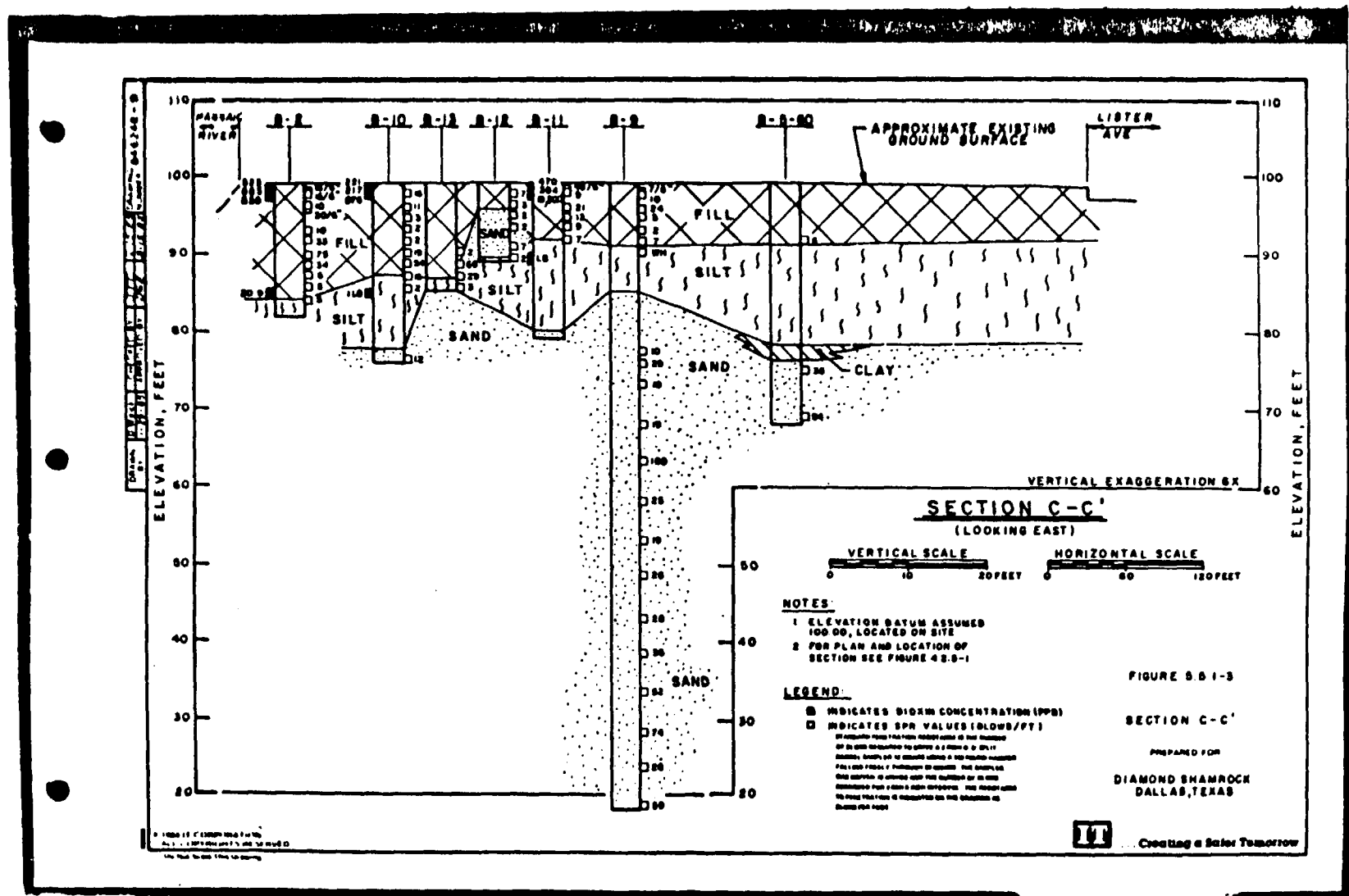
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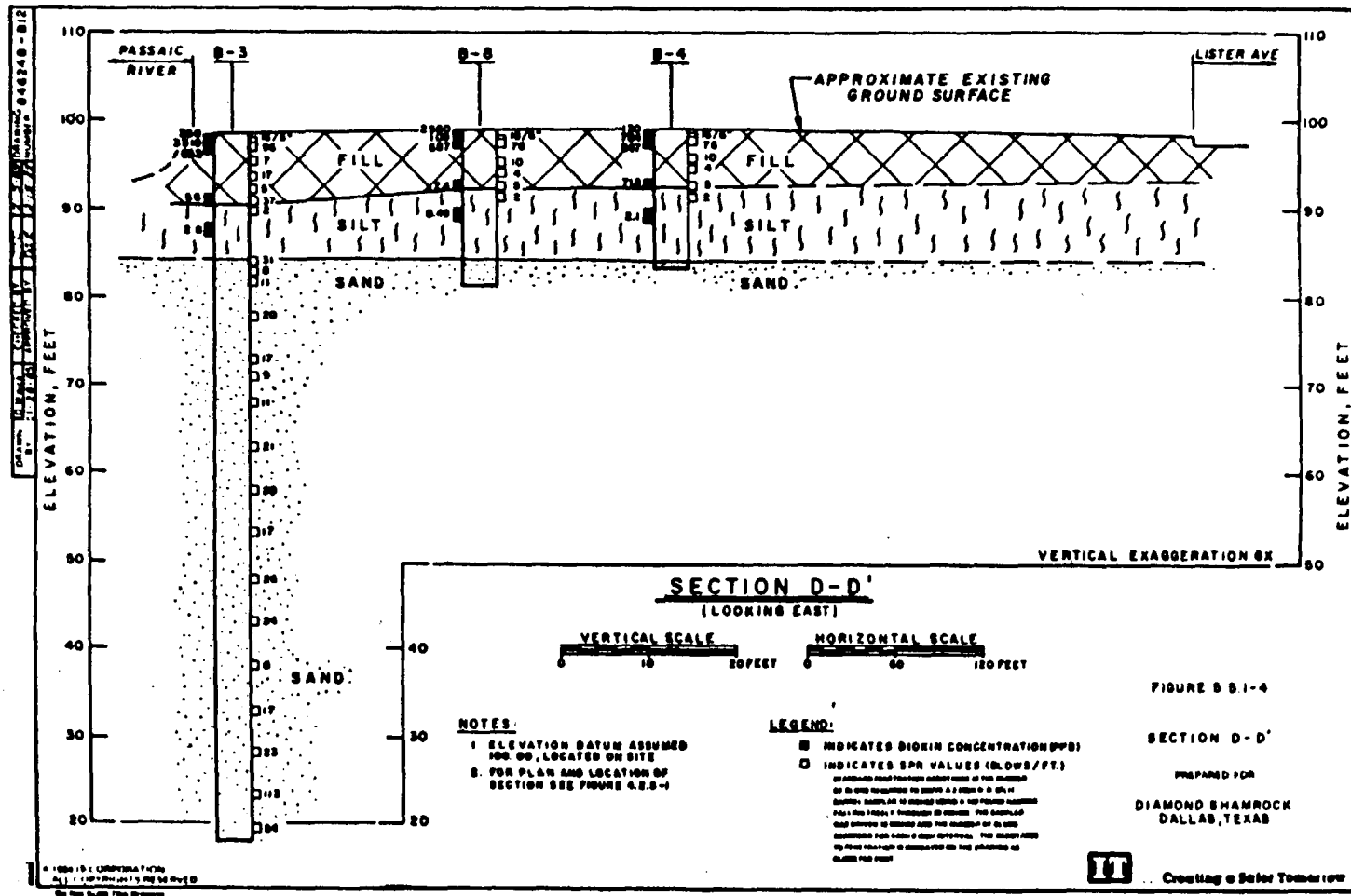




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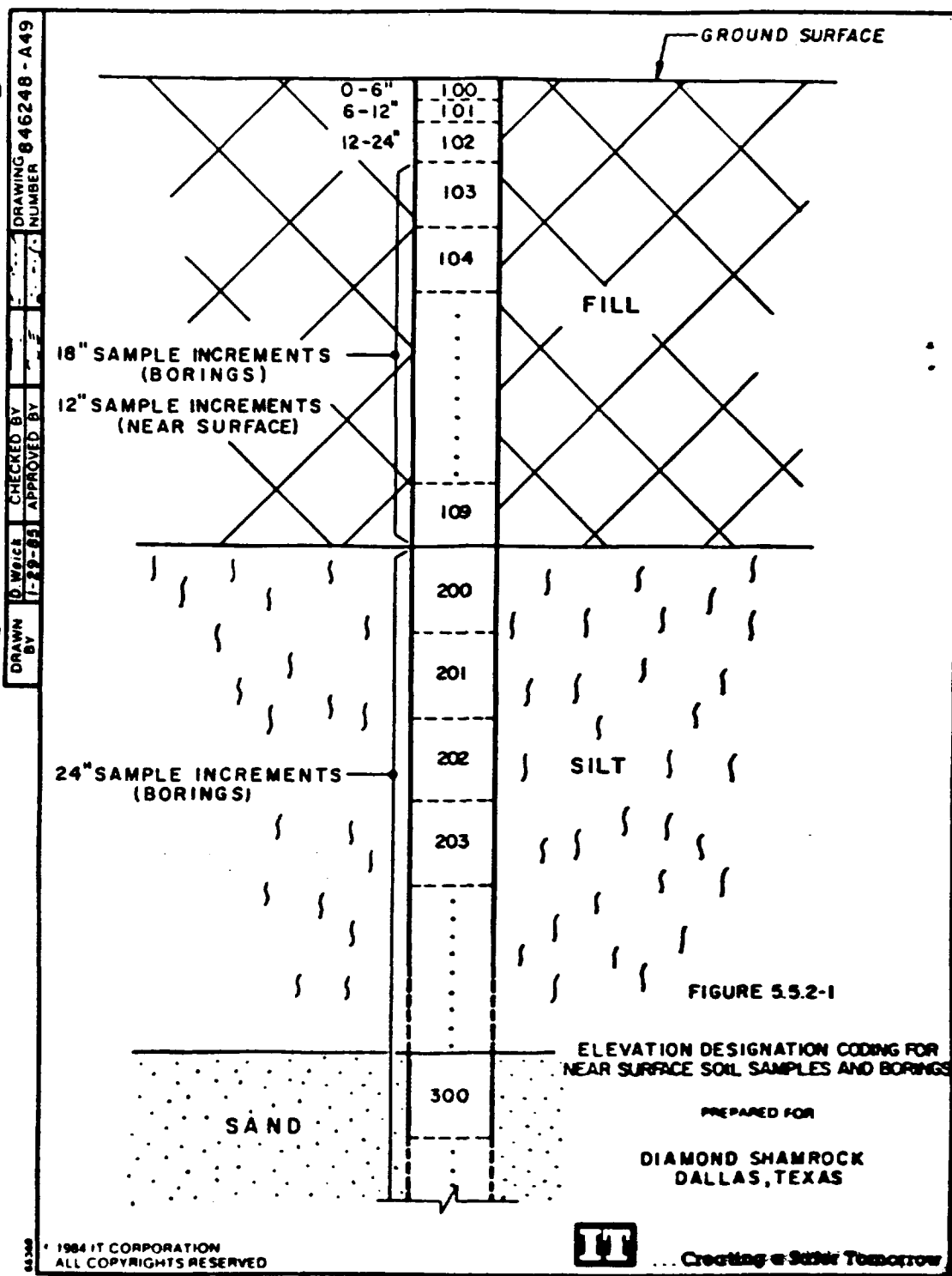


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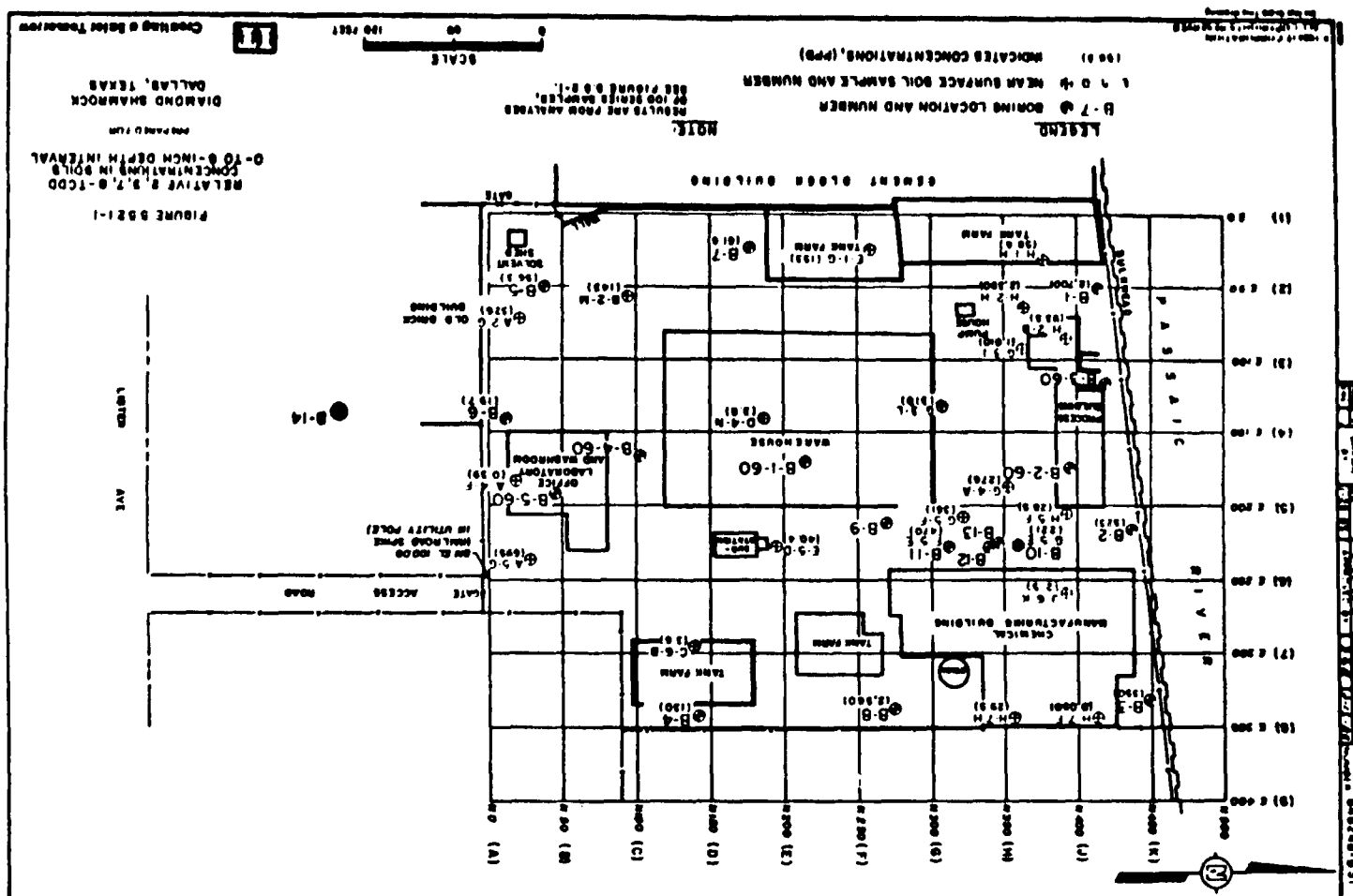
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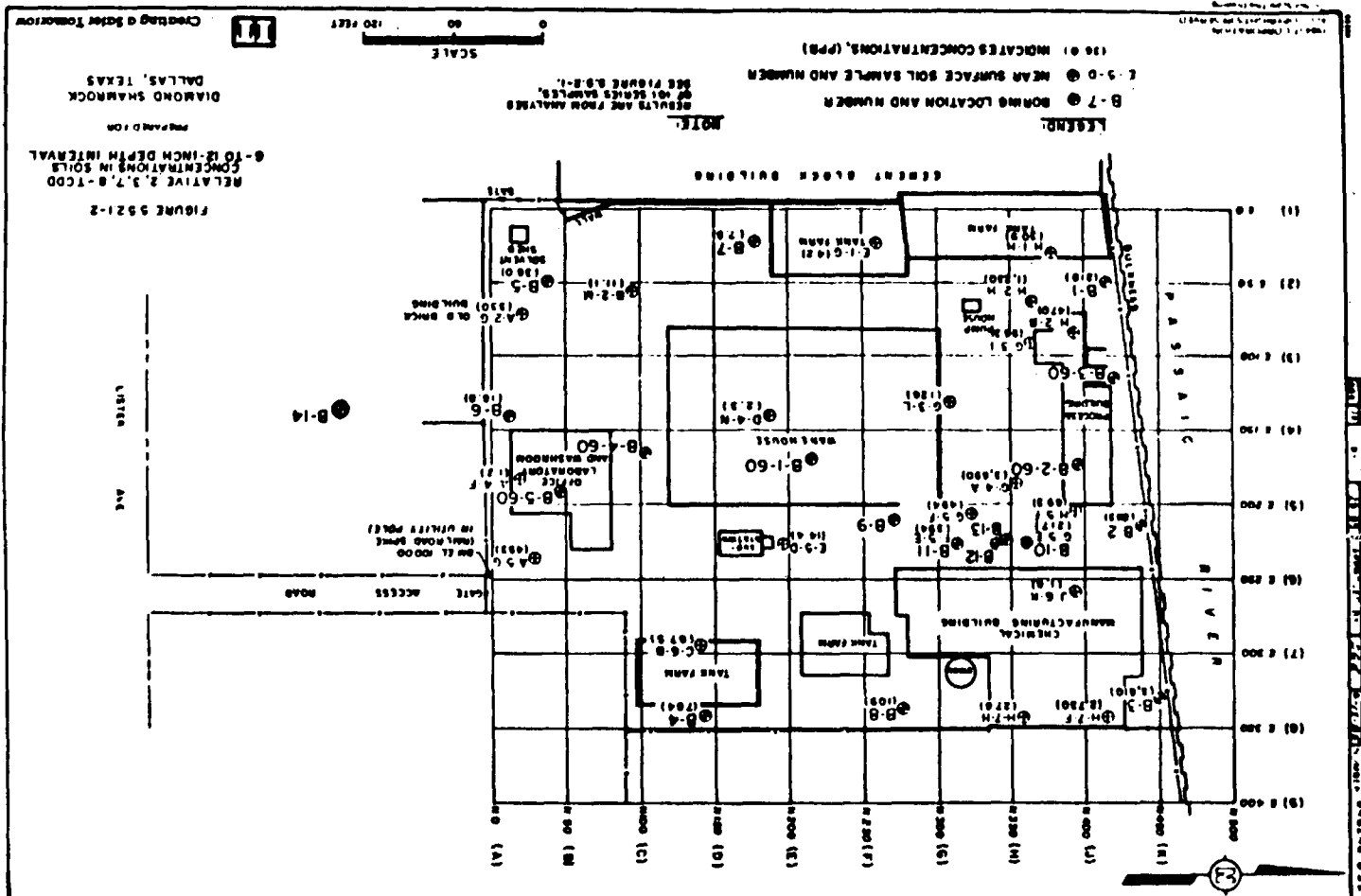
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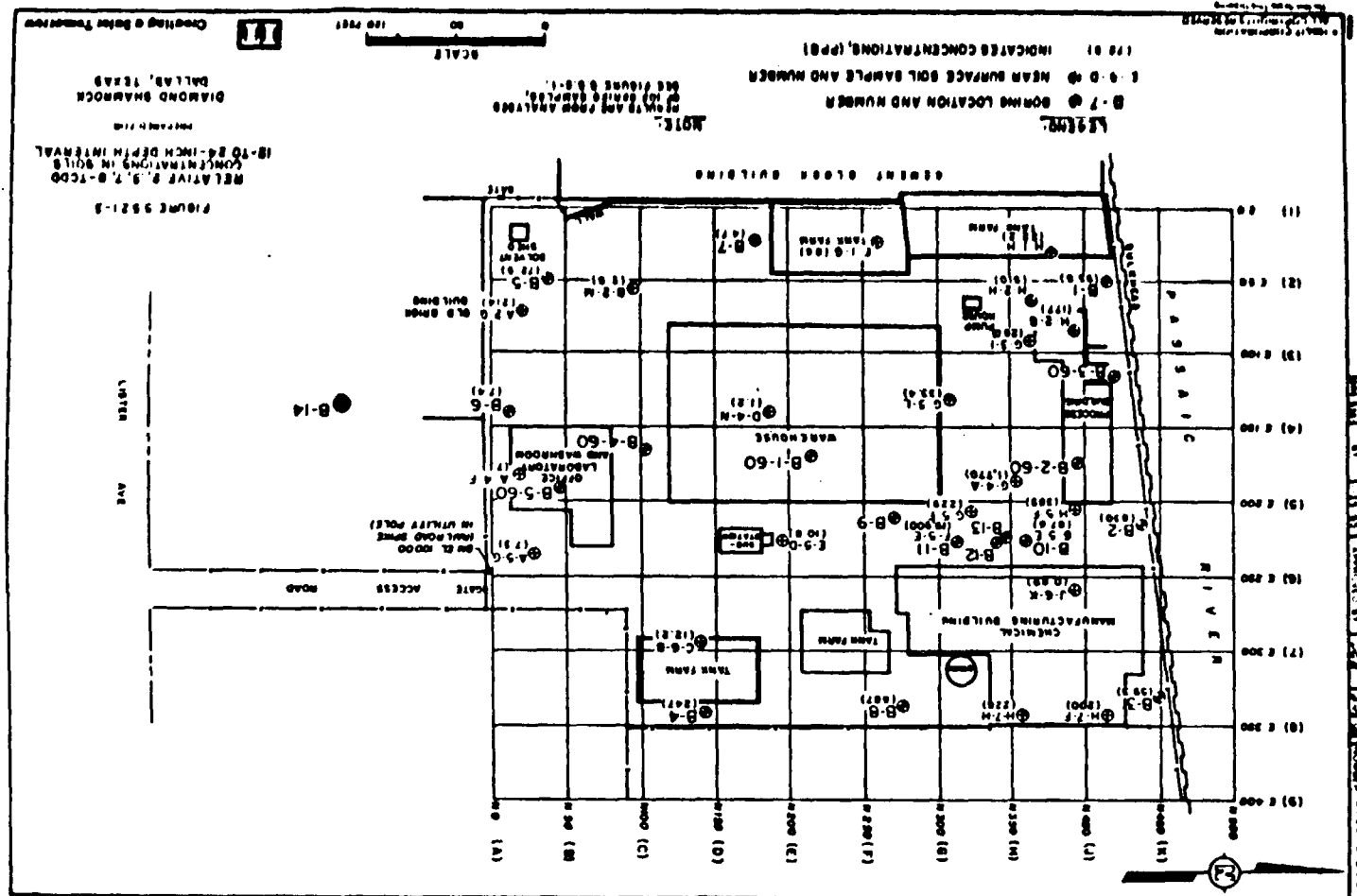
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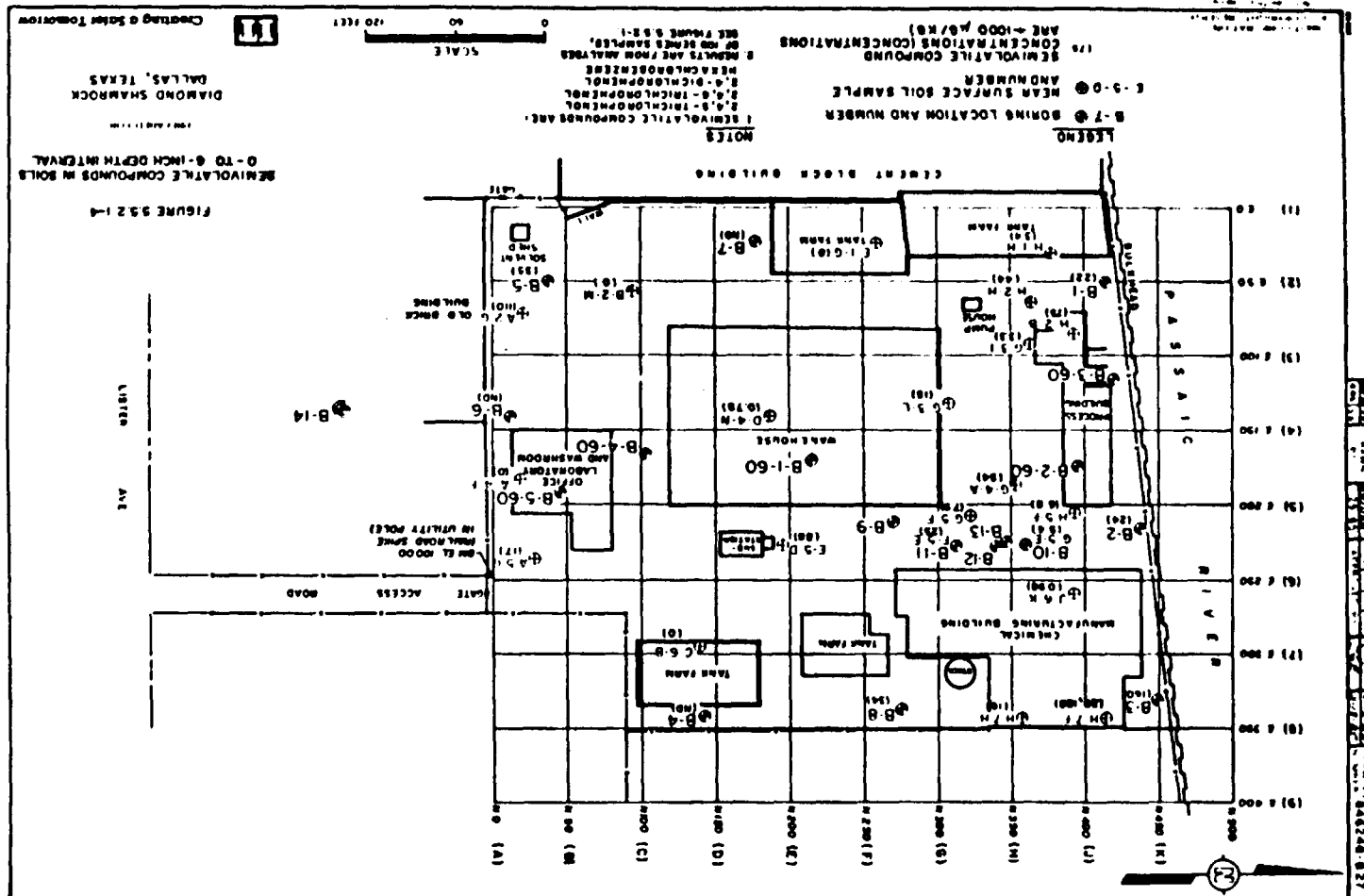
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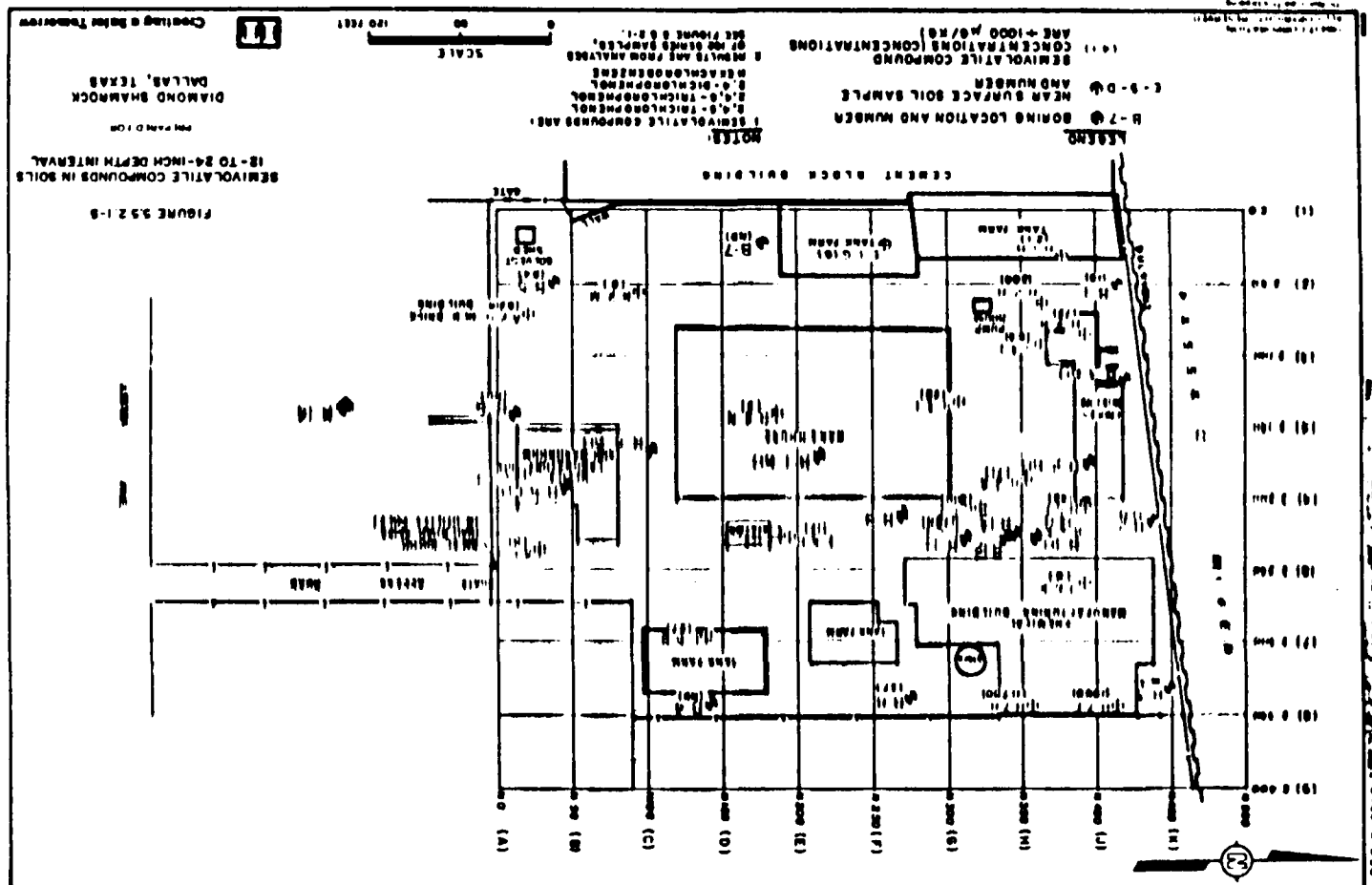
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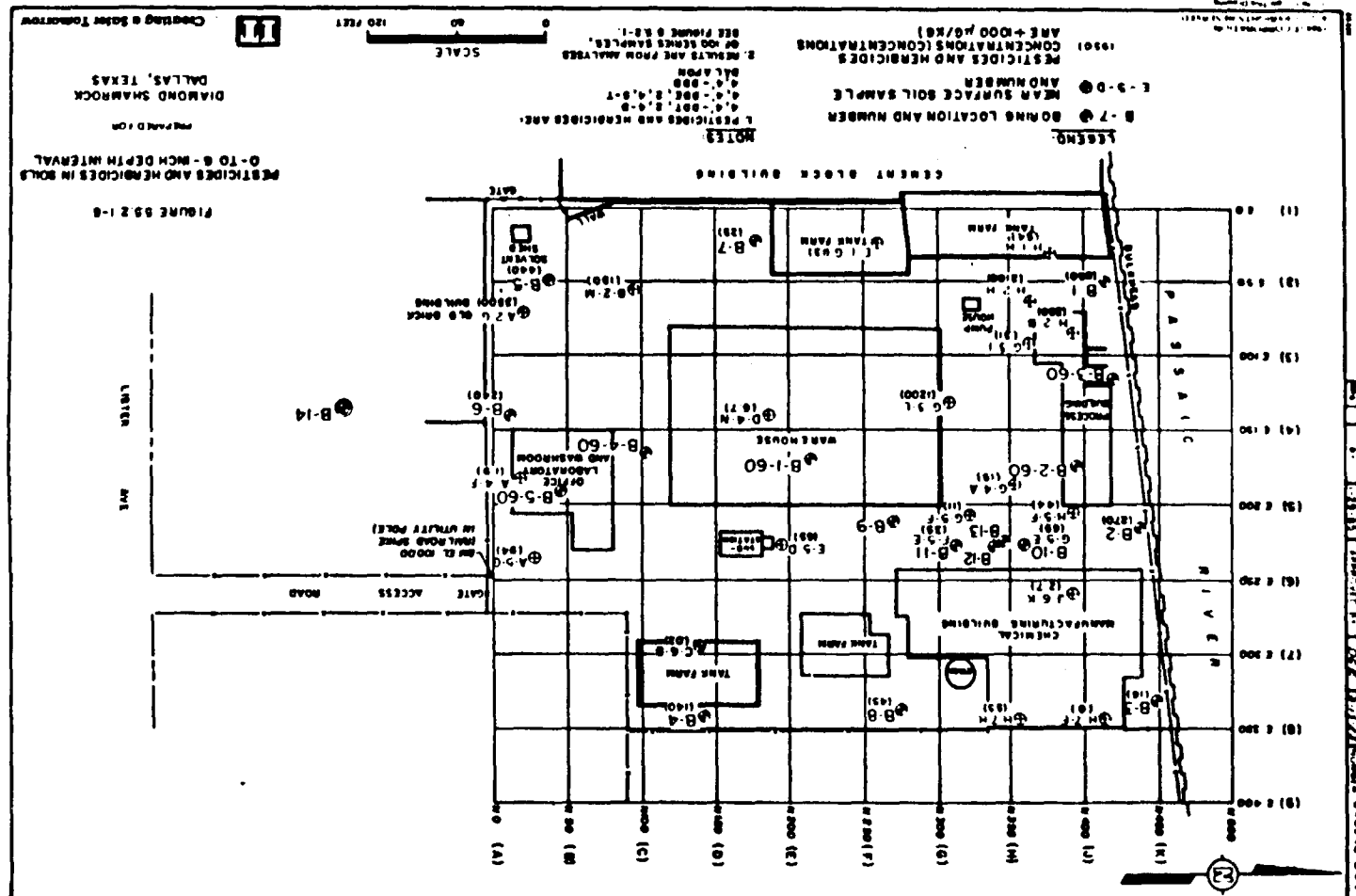
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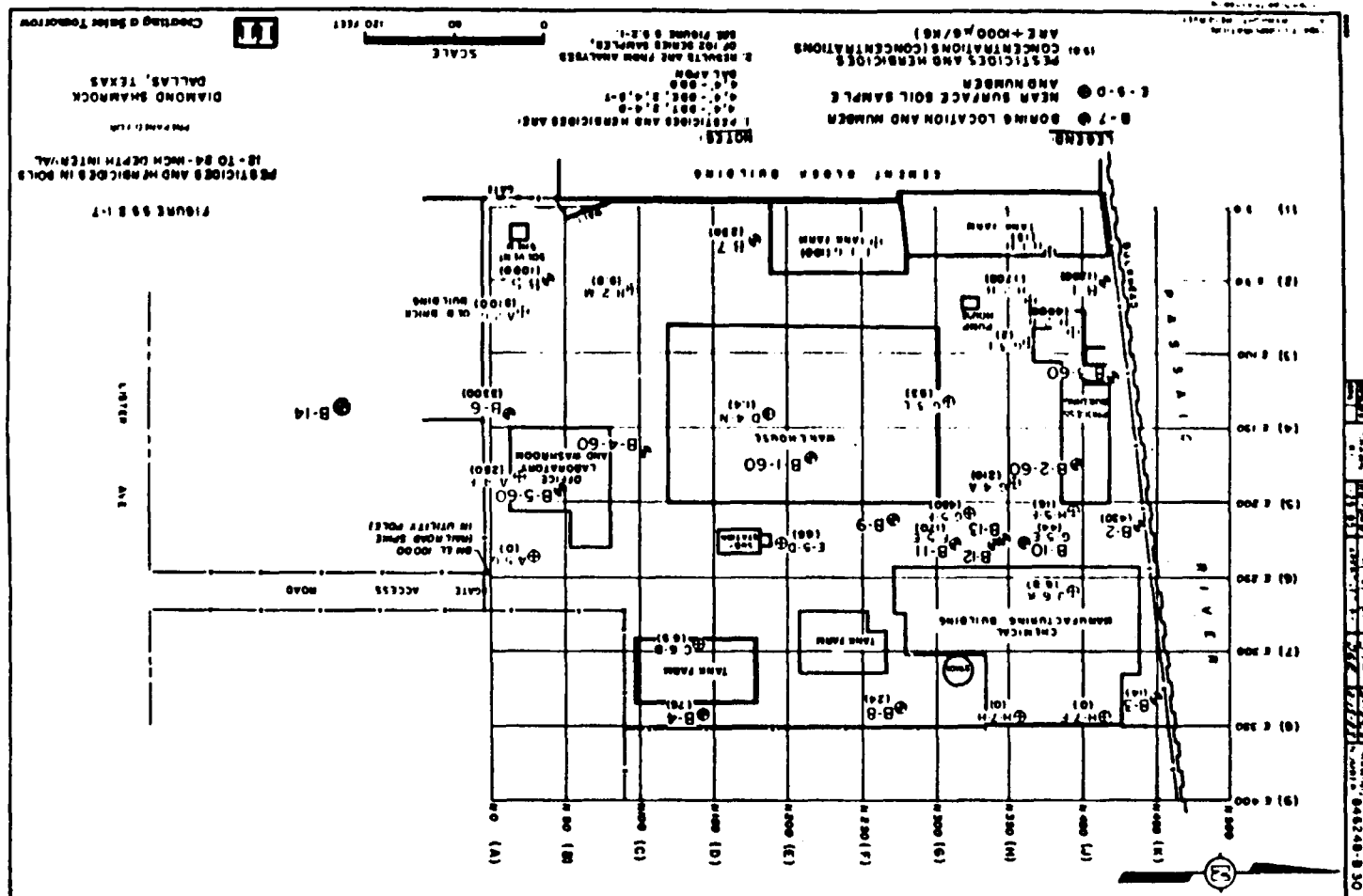
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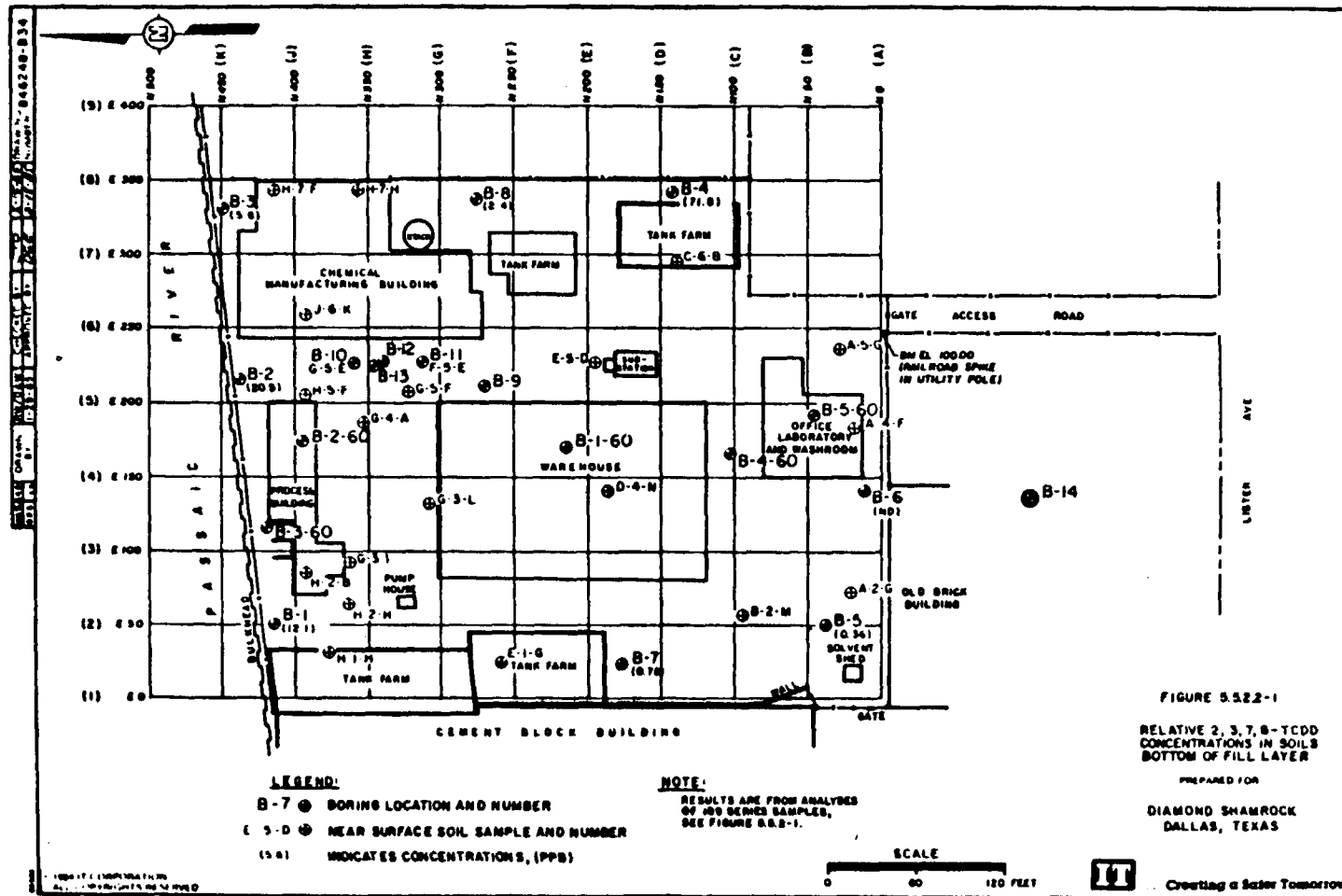
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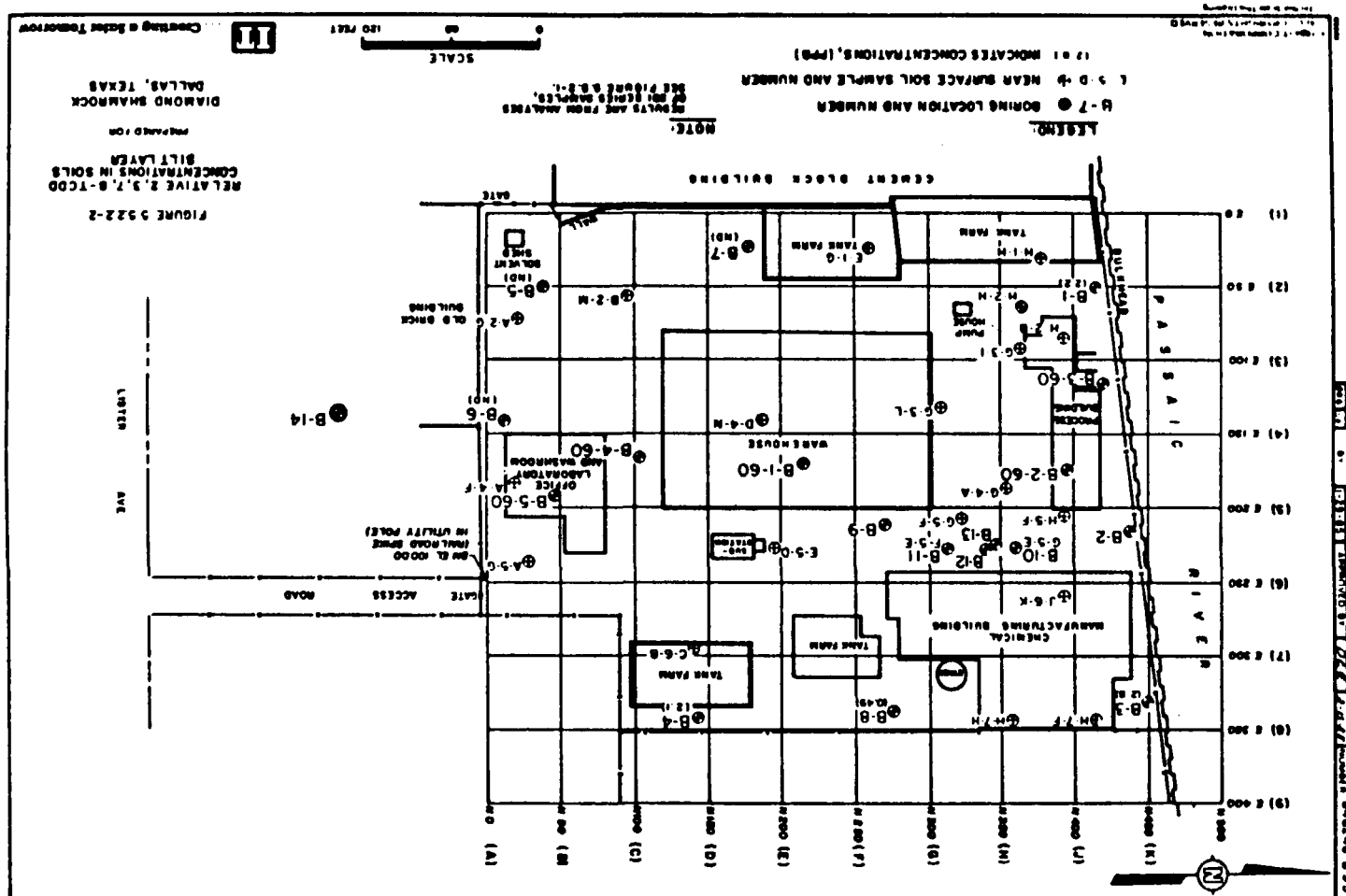


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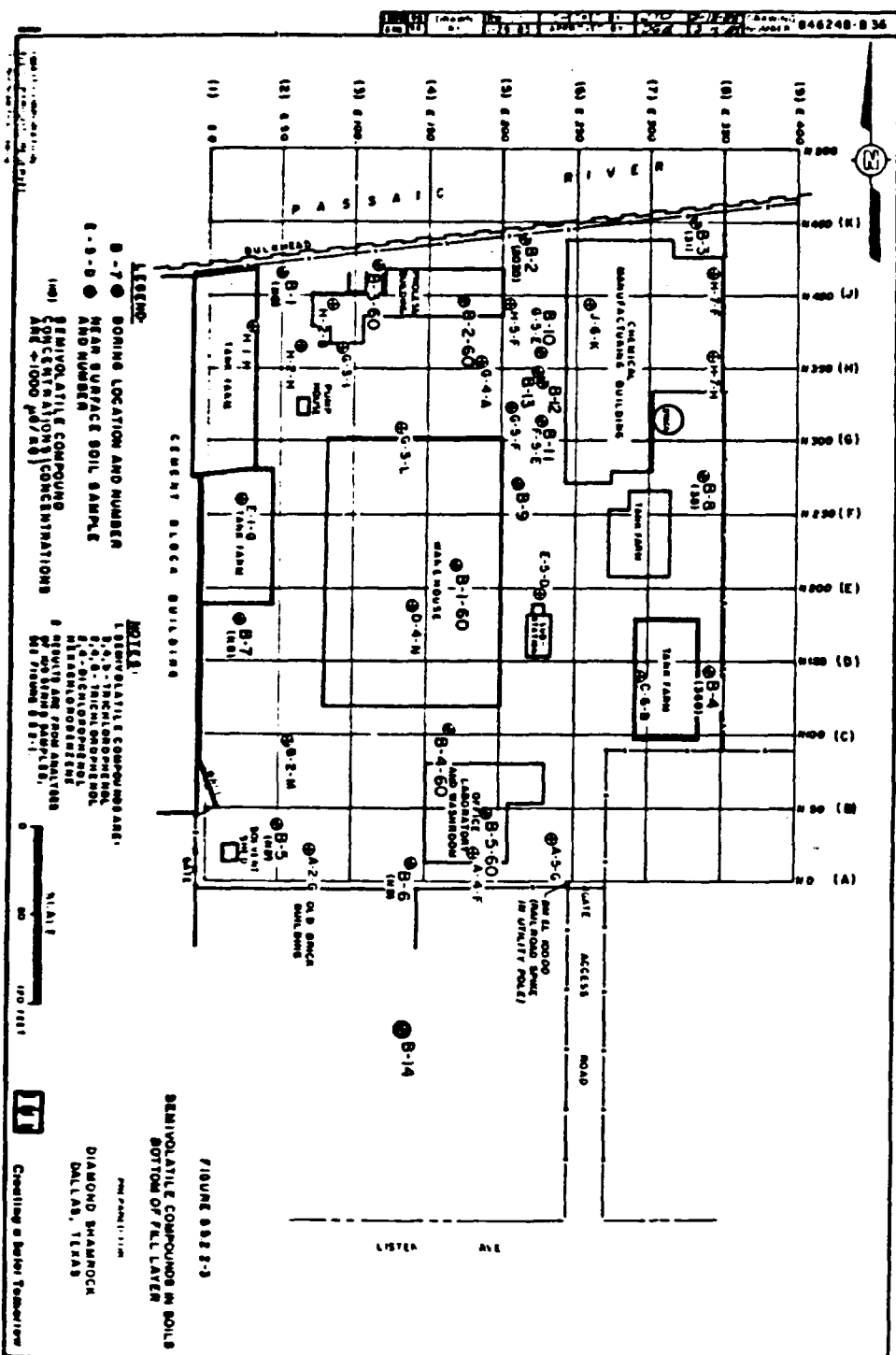
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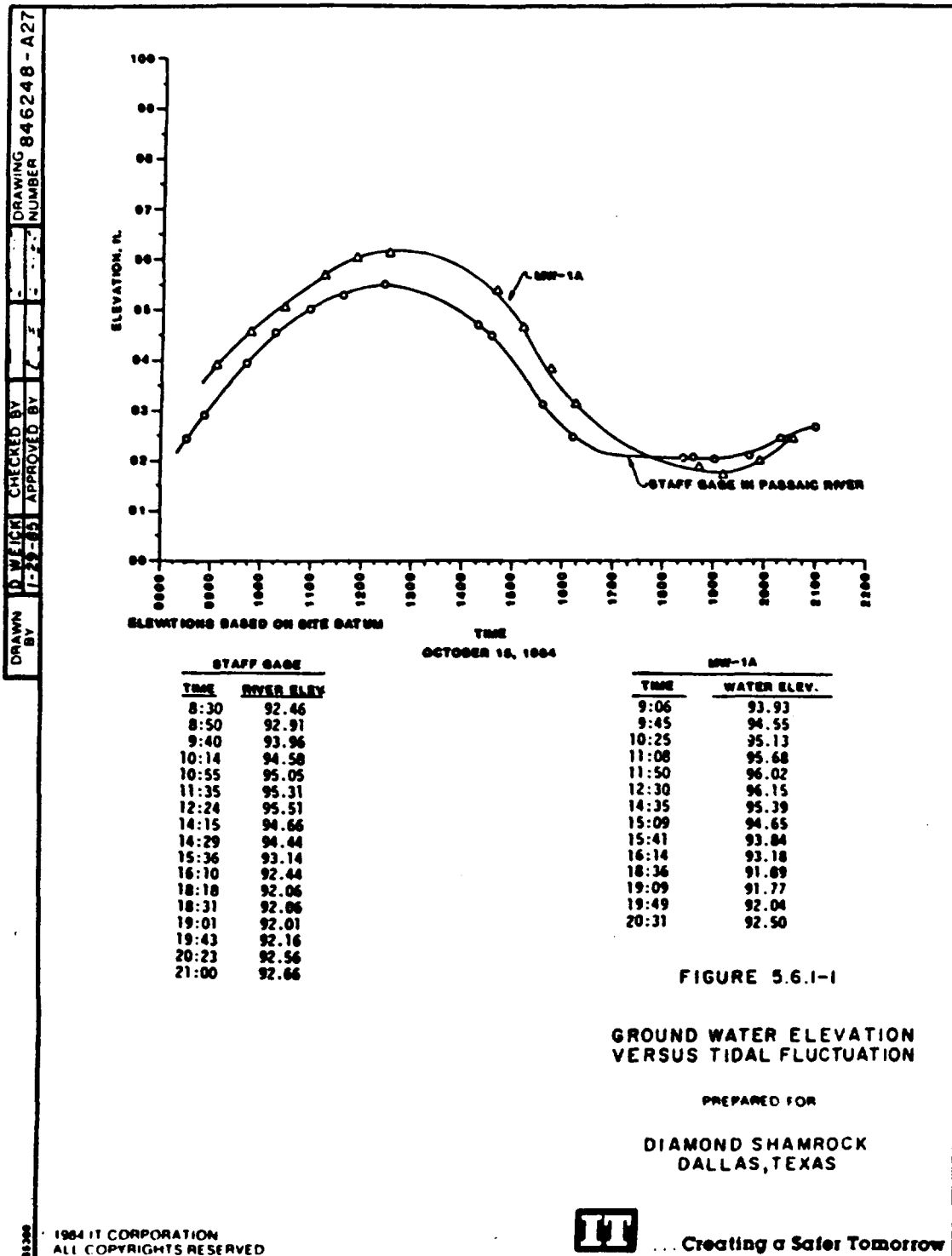
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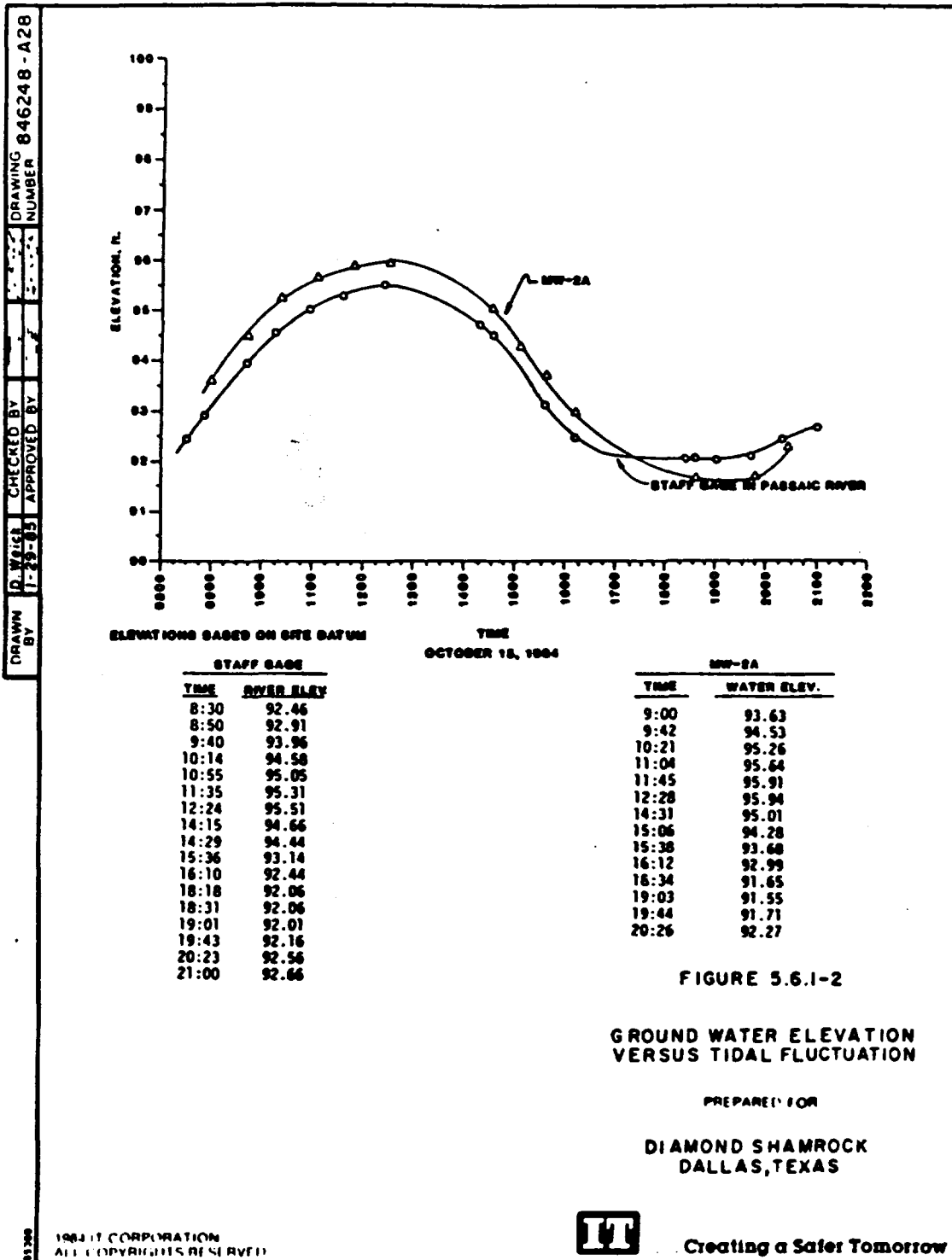




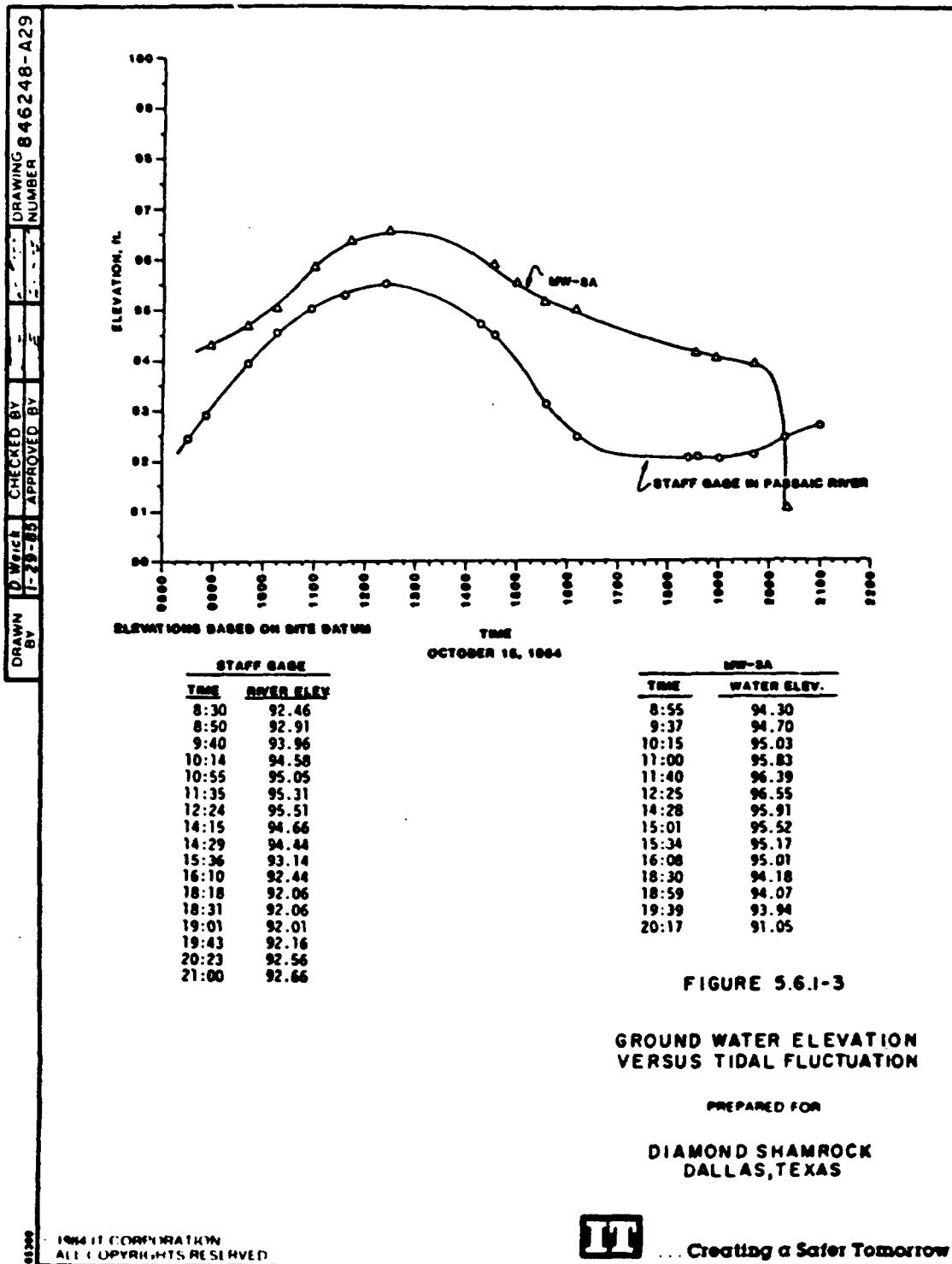
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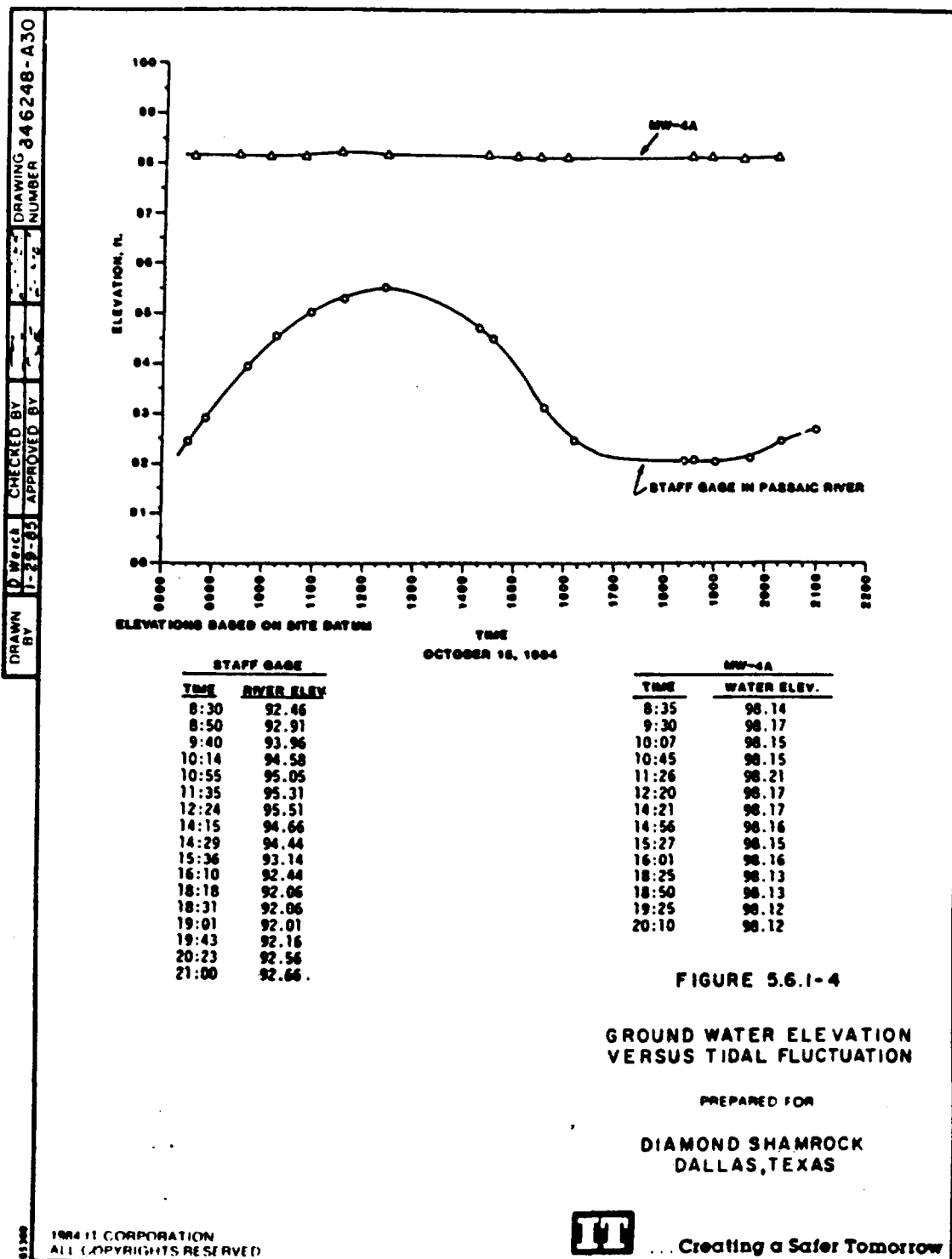


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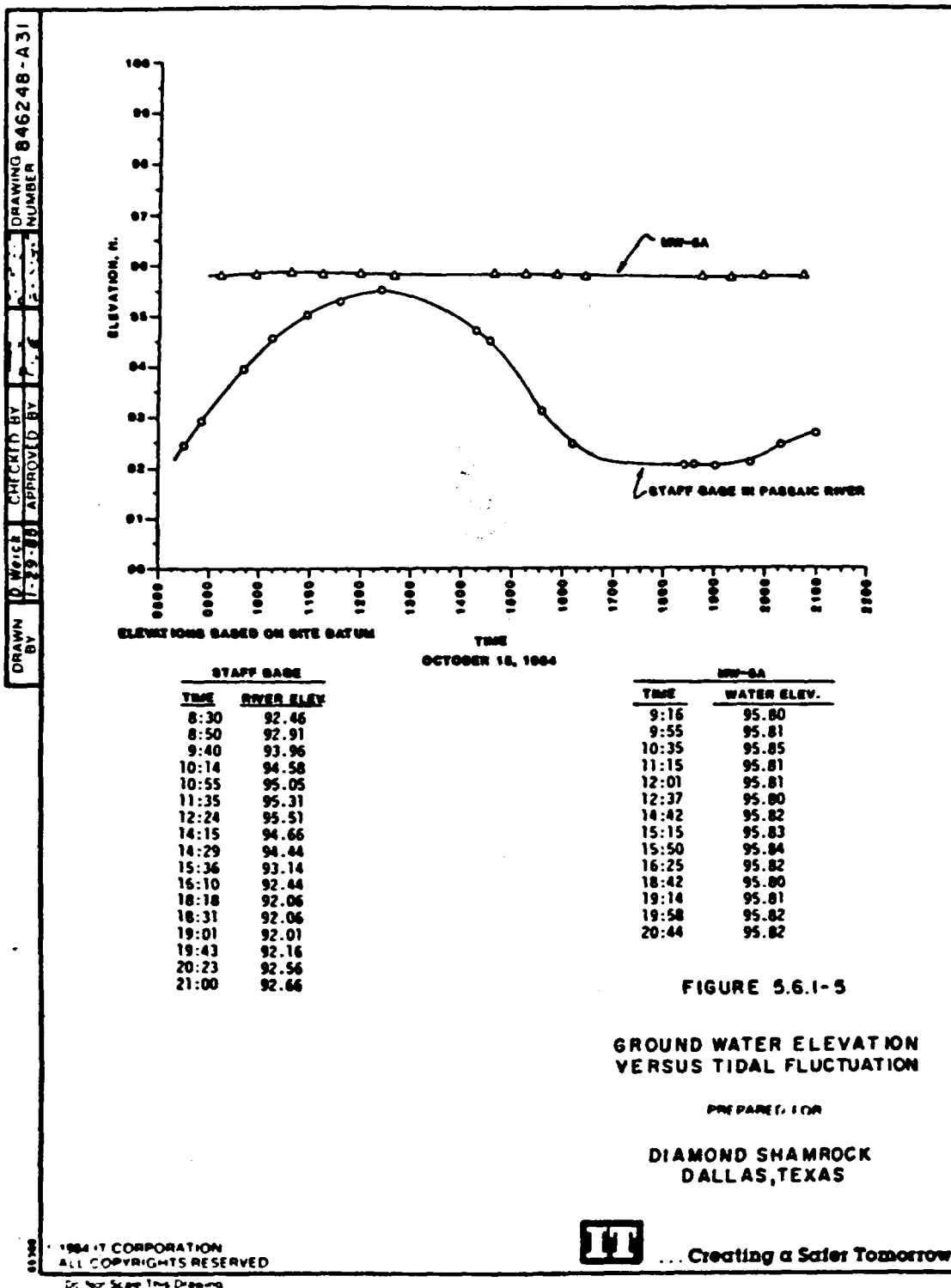
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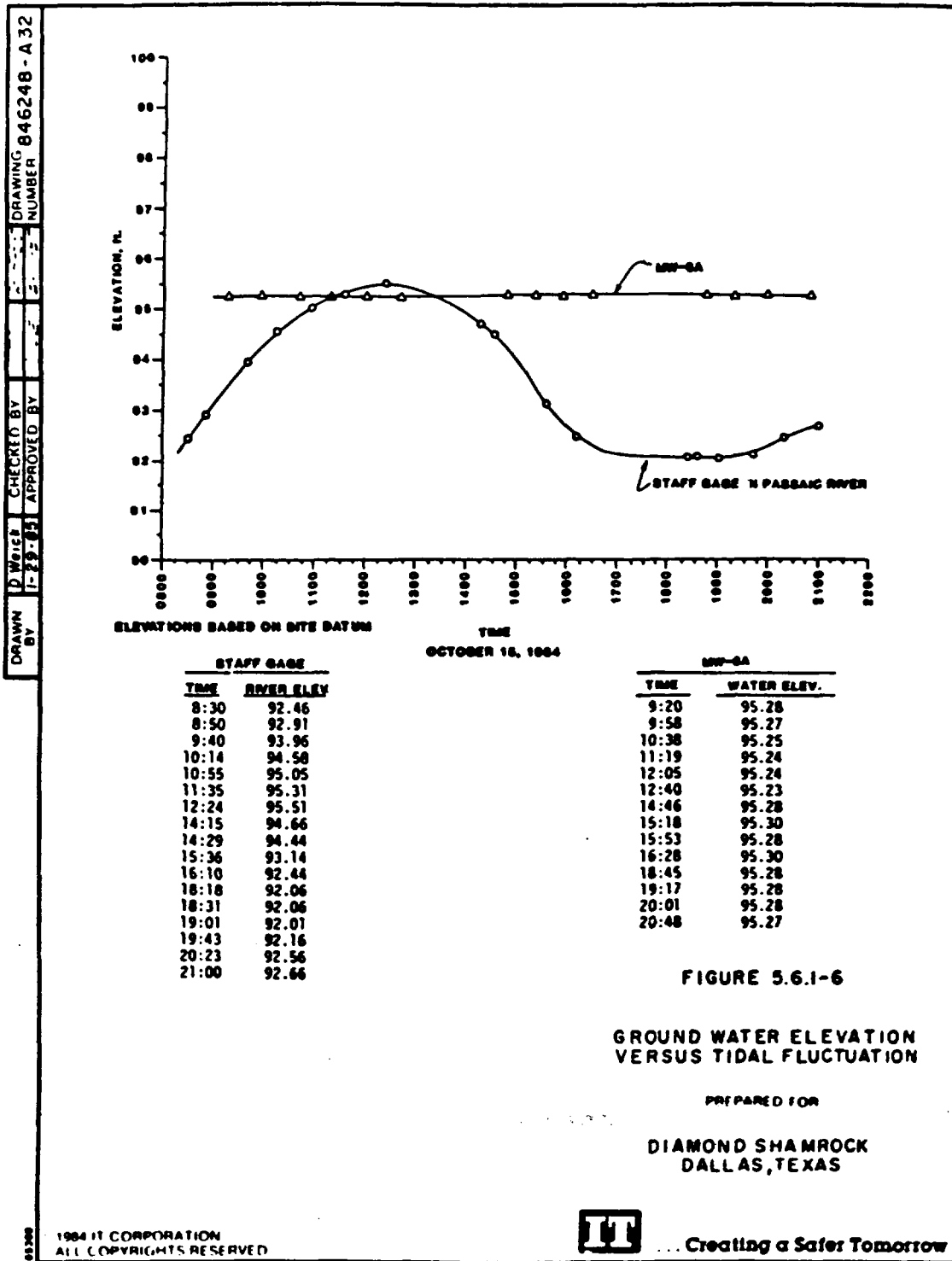


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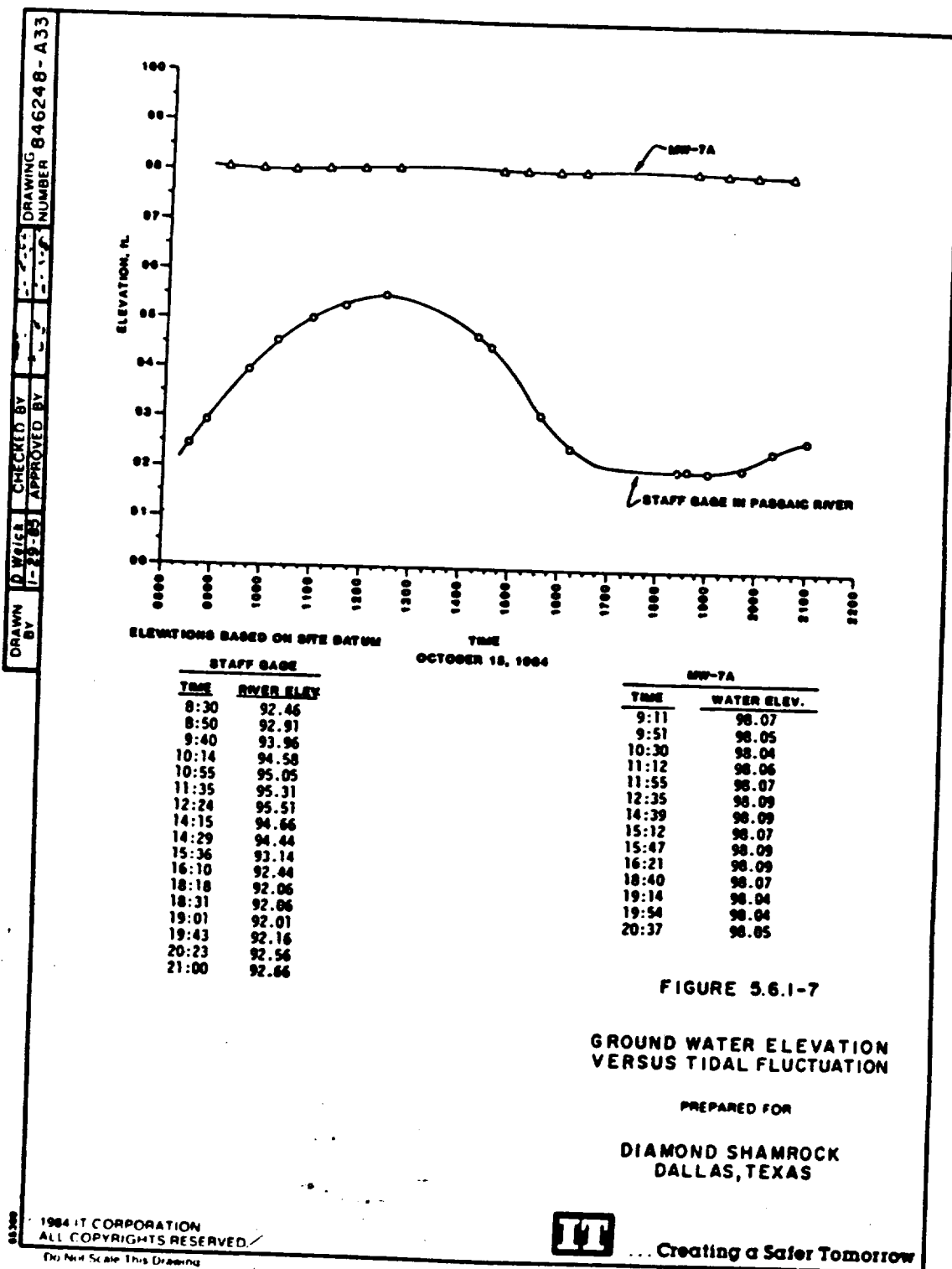




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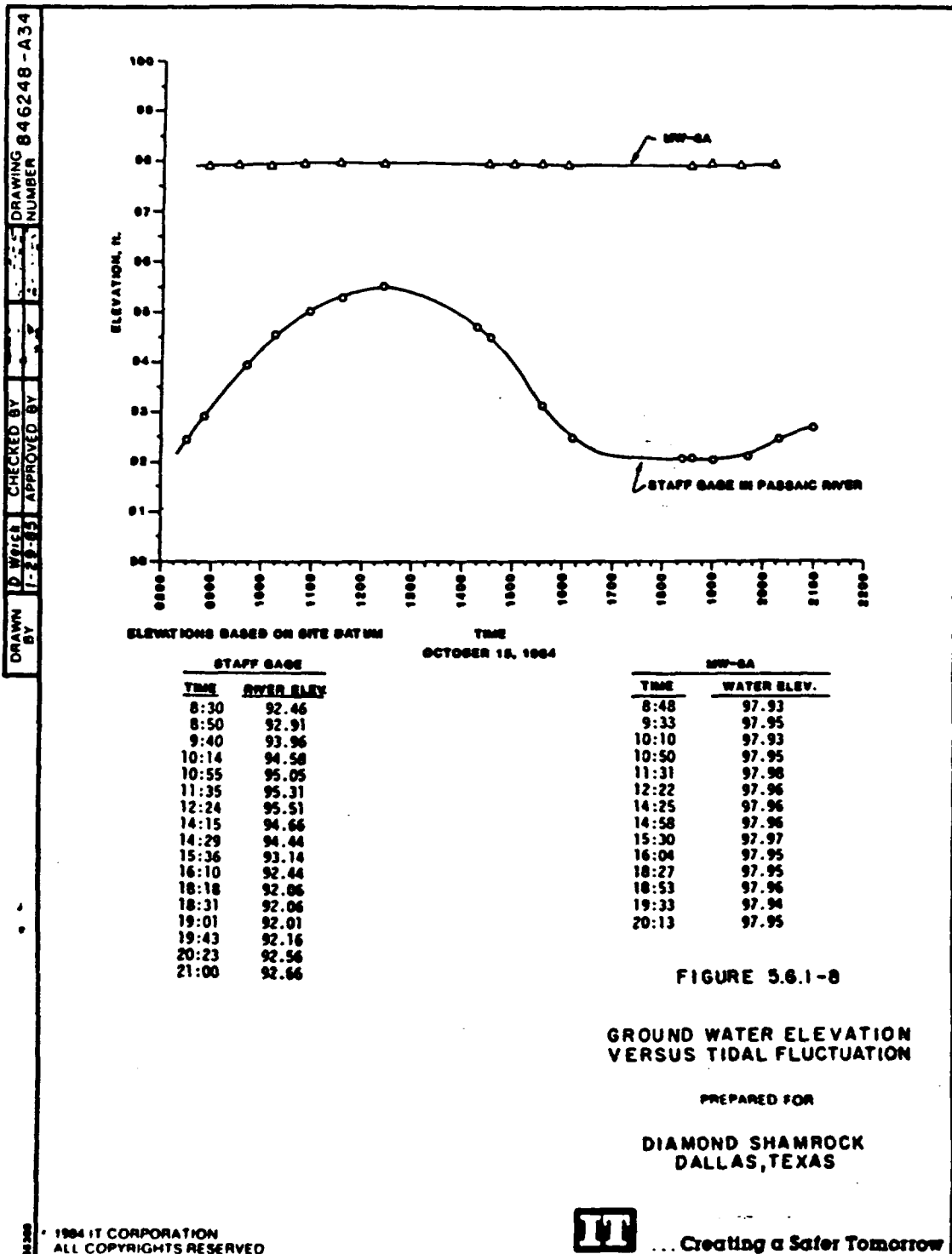
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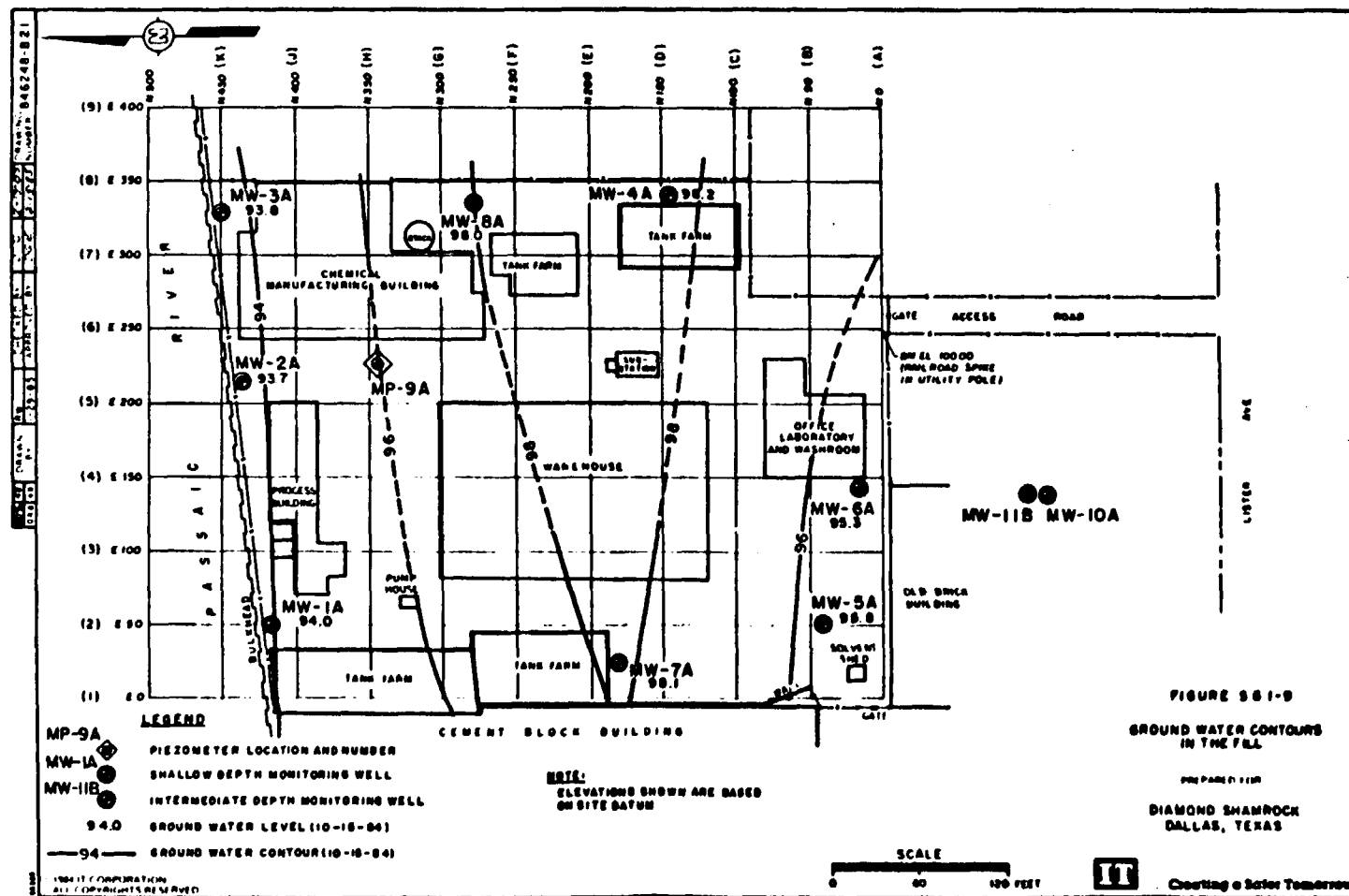
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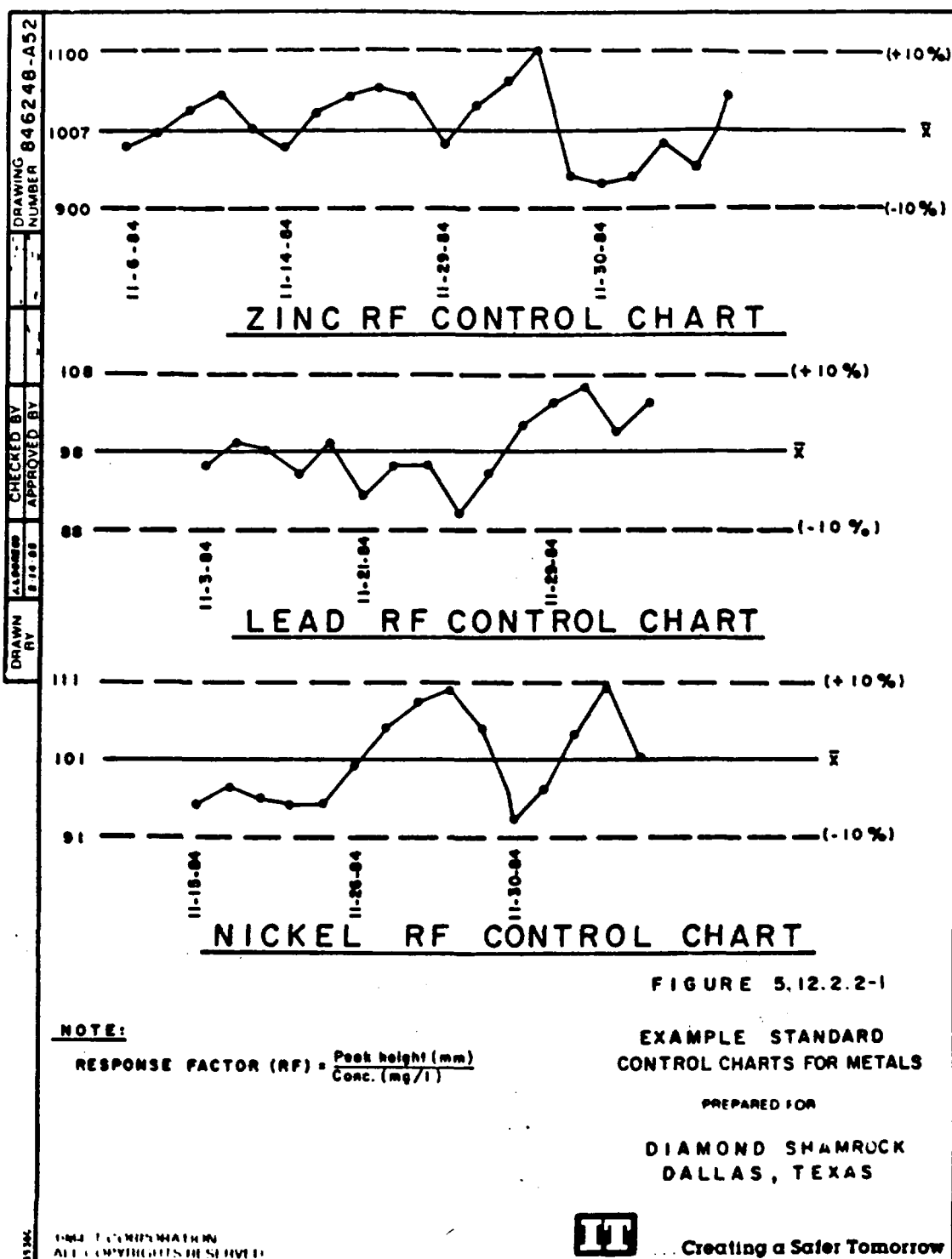
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6-1

6.0 SITE CHARACTERIZATION

6.1 AMBIENT AIR

The highest concentrations in the ambient air samples of total suspended particulates matter (TSP) and inhalable particulate matter (IPM) both occurred on September 25, 1984. The TSP concentration on that date was 254 milligram per cubic meter (mg/m^3) which exceeds the secondary 24-hour National Ambient Air Quality Standard (NAAQS) of $150 \text{ mg}/\text{m}^3$ for TSP, but is less than the 24-hour primary standard of $260 \text{ mg}/\text{m}^3$. The IPM concentration on September 25 was $196 \text{ mg}/\text{m}^3$ which is greater than the $150 \text{ mg}/\text{m}^3$ lower range for the proposed 24-hour IPM NAAQS.

The highest total metals concentration was $2.6 \text{ mg}/\text{m}^3$ and occurred on October 4, 1984. The relative concentrations of the various metals are consistent with those measured in Newark during the Airborne Toxic Elements and Organic Substances (ATEOS) studies. The most abundant metals were iron, lead, manganese and zinc, again paralleling the results of the ATEOS studies.

Dioxin was not detected on any of the days on which maximum concentrations of TSP, IPM, metals, VOC's, pesticides or PMA were measured. Dioxin was detected on September 10 and 24, at respective concentrations of 86 and 286 picograms per cubic meter (pg/m^3).

The highest total VOC concentration was $182 \text{ mg}/\text{m}^3$ which was measured on September 25. Major constituents were xylene, toluene and ethyl benzene, all aromatic hydrocarbons commonly emitted from petroleum refineries and plants manufacturing coatings and solvents; they are also constituents of aviation fuel. Other major constituents were trichloroethene and tetrachloroethene which are solvents widely used for dry cleaning and degreasing.

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Twenty-two drums representing 21 different lots (which comprised 499 of the 570 drums stored) were analyzed for dioxin. Fifteen of the analyses were positive, i.e., greater than one ppb, and seven produced non-detected results. The most significant dioxin contamination was detected in a drum containing solids and sludge labeled lot "Pit 3" (36 drums were included in this lot) at 8,750 ppb and from a drum containing yellow crystalline powder labeled lot "CQ" (11 drums were included in this lot) at 12,200 ppb. A weighted average of the lots determined to have positive dioxin concentrations indicates an estimated 230,000 pounds of material containing 940 ppb dioxin, or 98 grams. The 47 drums of "Pit 3" and "CQ" material contribute 93 grams of the total estimated quantity of dioxin.

In addition to the 570 drums already on the site, 125 drums of various waste were created during the site investigation undertaken by Diamond Shamrock. All of this material is currently stored in labeled drums in the warehouse. The location of each numbered drum is shown in Figure 6.2-1. The material stored includes soil, cement, and debris from drilling operations; water collected from the bailing of monitoring wells while sampling ground water; trash including bottles, polyethylene sheeting, boxes and paper; disposable items such as gloves, towels, and tyvek suits; and water and debris collected from the cleaning of decontamination water storage tanks and the diked areas around the tanks.

Material stored in drums 600 through 663 and 691 through 697 was collected during the site sampling activities. Drums 664 through 690 contain primarily water with some debris collected from inside the dike areas at the two major tank farms, and from the inside of Tank No. 2. Drums 698 through 715 were collected during the general site cleanup and background drilling operations on the Sherwin-Williams property. Drums 716 through 724 were filled during the final on-site cleanup activities.

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dust collected from flat elevated surfaces inside the building. The laboratory/office building is essentially the cleanest of the four major buildings with the highest levels detected in the floor of the basket room at 69.3 ppb and on a laboratory hood at 14,000 ng/m². The level detected in the laboratory hood is judged to be an isolated event because of the function of the hood. The stack and solvent shed had low levels of contamination both inside and outside of these structures. Contamination in the pump house, at 5.3 and 50.0 ppb, was notably higher.

In general, dioxin contamination is greater on the interior surfaces of all the buildings than on the exterior surfaces. Roofs have the lowest dioxin concentrations, while floors usually have the greatest. The laboratory/office building exterior walls had no positive dioxin results above ground level.

Due to the biased sample location strategy used, extrapolation of the results to quantify the total amount of dioxin in the structures is not feasible. However, an estimate of the volume of various types of materials in each of the major buildings that is potentially contaminated with dioxin has been made and is shown in Table 6.3-1. The total volume of material from all four major buildings is estimated to be 3,180 cubic yards.

Associated with the process and chemical manufacturing buildings are a number of sumps and the industrial sewer system.

All of the sumps inspected during the site investigation had been previously cleaned or emptied to some degree. However, enough residual material remained in each to enable the collection of bulk samples. A total of eight sump samples were taken, three from within the chemical manufacturing building, two outside the chemical manufacturing building, and three just outside the process building. Four sewer samples were

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The thickness of the fill ranges from eight feet along the southern property boundary to 15 feet in the northwest corner of the property. Along the western boundary of the site, the old river channel is fairly well defined. The fill is approximately eight feet thick up to the point where the old river channel is encountered north of Boring B-7. In the central portion of the site, the edge of the old channel is defined just north of Boring B-11. The fill has a fairly uniform thickness of eight feet to a point just north of Boring B-11 and gradually increases in thickness in the direction of the bulkhead at the northern site boundary. It is 15 feet in thickness at Boring B-2. Along the eastern boundary of the site, the old river channel has not significantly intruded into the property. At the southeastern corner of the site, the fill is six feet in thickness. The fill increases only slightly to a thickness of eight feet at the bulkhead in the northeastern corner of the site.

Subsection 5.5.2.1 describes the dioxin levels detected in the fill layer resulting from the analysis of near-surface soil samples and shallow depth boring soil samples. The samples obtained in the fill were analyzed for dioxin in increments of depth from zero to 6, 6 to 12, and 12 to 24 inches. Soil samples obtained from the borings at depths greater than 24 inches were archived for possible future analysis with the exception of several samples (109's) obtained immediately above the silt layer.

For the zero- to 6-inch depth, dioxin concentrations varied from 0.39 ppb to 9,050 ppb. For 6- to 12-inch depths, the dioxin concentration varied from 1.2 ppb to 3,690 ppb. The analysis of samples obtained in the 12- to 24-inch depth interval resulted in concentrations ranging from 0.92 ppb to 19,500 ppb. Table 6.4.2-1 summarizes dioxin concentration ranges for the three depth intervals. All samples to the 24-inch depth analyzed for dioxin had positive results.

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feet across the site with no identifiable trends in deposition. The organic silt layer is composed of decayed grasses rooted in organic material produced from earlier growth. It has decayed to the present deposit of fibrous, highly organic peat with intermingled silt and humus.

The lowermost layer of the silt deposit extends across the site and is probably the depositional result of tidal flat processes. It is a fine-grained sediment deposited in a low-energy environment. Intermingled in this layer are some roots and thin sand and gravel layers. The contact between the two silt units and the underlying glacio-fluvial sands is generally gradational. The thickness of this lower unit varies from three to eight feet, with the thickest deposit detected in the off-site (Sherwin-Williams) wells.

Initially, seven samples in the silt layer were analyzed for dioxin. Subsequently, two additional silt samples, from B-10 and B-11, were also analyzed. It was not expected that dioxin would be detected, but six of these nine samples indicated results above the method detection level, ranging from 0.49 to 11.8 ppb dioxin. Because of these positive results, five additional silt layer samples were selected from the archived Shelby tubes and analyzed. The five archive samples resulted in two results above detection limits. The highest dioxin result in any silt sample was 11.8 ppb from Boring B-10, which is adjacent to the chemical manufacturing building. It is noted that the upper organic silt layer is not present in this area. The remaining positive values ranged from 1.2 ppb to 2.8 ppb dioxin.

Analysis for priority pollutants was not performed on silt layer samples.

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in thickness from 3 to 11 feet. Data from site borings indicate that this silt layer is probably continuous across the site.

The fill material on the site constitutes a "surficial aquifer" and the silt layer underlying the fill has a lower permeability, restricting the downward movement of ground water contained in the fill. Table 5.6.1-1 presents a summary of monitoring well data for the fill material. The last column of the table shows that the saturated thickness of the permeable zones in the fill ranges from about two to eight feet. The saturated thickness for monitoring wells MW-1A, 2A, and 3A is likely to range both higher and lower than the values shown in Table 5.6.1-1 because of the effect of tidal fluctuations on surficial ground water levels near the river. Although fill layer is denoted as the surface aquifer, it is very limited in extent and does not serve as a source of potable or industrial water usage.

The glacio-fluvial sand deposit underlying the silt layer is a significant water-bearing unit. The three deepest borings on site (B-1, B-3, and B-9, all drilled to 81.5 feet) indicated this unit to be at least 62 to 68 feet thick at the boring locations. The unit appears to be completely saturated. Some possible industrial usage of water from this unit is indicated in Table 3.3.2-1. It is not known whether a glacio-lacustrine clay layer separates the glacio-fluvial sand deposit and the shale or sandstone bedrock at the site as it does in many parts of the area (Nichols, 1968). The glacio-lacustrine layer could retard downward flow in the sand unit.

Bedrock is composed of Triassic sandstones and shales of the Brunswick Formation. Based on well data from nearby areas, it appears that the Brunswick Formation beneath the site is fractured. The Brunswick aquifer is the source of potable water (Table 3.3.2-1) in the surrounding area.

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borings, it is estimated that the hydraulic conductivity of the glacio-fluvial deposit ranges from 10 to 50 feet per day. This is the same range as the surficial fill.

Based upon comparison with values reported for similar materials in the literature, it is estimated that 0.003 feet per day (1.0×10^{-6} centimeter per second) is a reasonable value for the average vertical hydraulic conductivity of the silt. This is a low value and indicates a significant potential for retarding the flow of ground water downward from the fill.

6.5.4 Ground Water Flow

Based upon the ground water level measurements and slug tests performed in the eight monitoring wells, estimates of ground water flow directions and rates in the fill were made. Estimates of the vertical downflow of ground water from the fill through the silt to the sand can also be made. Because only one monitoring well was installed in the glacio-fluvial sand and none was installed in the Brunswick Formation, flow directions and rates can not be determined for these units.

Ground Water Flow in the Fill

Mean ground water level elevations and approximate ground water level contours for the fill at the site are shown in Figure 5.6-9. The data were obtained on October 14, 1984. Mean values, defined as the average of the maximum and minimum values in a tidal cycle, are shown for monitoring wells MW-1A, 2A, and 3A (obtained from Figures 5.6-1, 5.6-2, and 5.6-3, respectively). These mean values were determined on October 15; the corresponding water levels for the other monitoring wells were observed on the same day.

Ground water flow velocities in the surficial fill at the site were computed from the gradients (piezometric head divided by distance) developed from Figure 5.6-9 and hydraulic conductivities presented in

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Ground Water Flow in the Lower Units

One monitoring well was installed in the glacio-fluvial sand, but hydraulic conductivity testing was not performed. Therefore, insufficient information is available to estimate flow rate and direction for this unit. Similarly, no information regarding the Brunswick Formation was obtained as part of this investigation, since none of the borings reached this unit.

6.5.5 Extent of Contamination

Analysis of samples taken from the eight on-site wells has confirmed the presence of dioxin in the ground water. Contamination was present in 15 of 17 water samples collected, ranging from 0.0059 ppb in monitoring well MW-5A to 10.4 ppb in MW-2A.

The results indicate that contamination is greatest at the north end of the site along the river, near the process and chemical manufacturing buildings. Monitoring wells MW-5A, 6A, and 7A, located at the south and southwestern portions of the site, consistently had the lowest dioxin levels, ranging from ND to a maximum of 0.016 ppb; monitoring wells MW-4A and MW-8A along the eastern edge at the site had dioxin levels ranging from 0.20 to 1.1 ppb; monitoring wells MW-1A, 2A, and 3A located on the northern edge of the site showed levels ranging from 0.03 to 10.4 ppb dioxin.

Three ground water samples from monitoring well MW-2A in the center of the north end of the site had dioxin results of 7.9, 4.3 and 10.4 ppb, all significantly higher than any other ground water sample dioxin result. These very high levels of dioxin in water are probably the result of the presence of a contaminated solvent or other carrier that has solubilized in the ground water or because dioxin-contaminated colloidal soil particles are suspended in the water.

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A total of 15 sediment samples were analyzed for priority pollutants--10 at depths of zero to 12 inches and five at depths of 12 to 24 inches. Subsequent analysis confirmed the presence of a number of compounds.

In general, the upper (zero to 12 inch) samples showed higher values of semivolatile organics, equivalent values of inorganics and volatile organics, and lower values of pesticides than the 12- to 24-inch depth interval samples. The compounds with the highest concentrations detected were chemicals produced on the site, including 2,4-DCP, 2,4,5-TCP and their esters. Other compounds found in a significant number of the samples included the various metals, fluoranthene, phthalates and pyrene.

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TABLES

TABLE 6.3-1
ESTIMATED VOLUME OF CONTAMINATED MATERIALS
FROM MAJOR BUILDINGS

BUILDING	(SLAB) CONCRETE (yd ³)	CONCRETE BLOCK OR BRICK (yd ³)	WOOD OR PLASTER (yd ³)	ROOF MATERIAL (yd ³)	STEEL GRATING (yd ³)	STRUCTURAL STEEL (yd ³)	CORRUGATED ASBESTOS PANELS (yd ³)	TOTAL (yd ³)
Office and Laboratory	100	310	180	110	-	-	-	700
Warehouse	480	160	40	-	-	100	30	810
Process	110	110	-	25	40	100	15	400
Chemical Manufacturing	<u>475</u>	<u>610</u>	<u>-</u>	<u>25</u>	<u>60</u>	<u>80</u>	<u>20</u>	<u>1,270</u>
TOTAL	1,165	1,190	220	160	100	280	65	3,180

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TABLE 6.4.2-1
2,3,7,8-TCDD CONCENTRATION RANGES
VERSUS NUMBER OF SAMPLES ANALYZED

DIOXIN CONCENTRATION RANGE (ppb)	FILL LAYER DEPTH INCREMENT (inches)		
	0 to 6	6 to 12	12 to 24
<50	7N* + 1B**	9N + 3B	11N + 2B
50 to 200	4N + 3B	4N + 1B	3N + 3B
200 to 500	6N + 1B	5N + 1B	4N + 1B
>500	4N + 3B	3N + 3B	3N + 2B

*N refers to samples collected in the near surface soil sampling program.

**B refers to samples collected in the boring soil sampling program.

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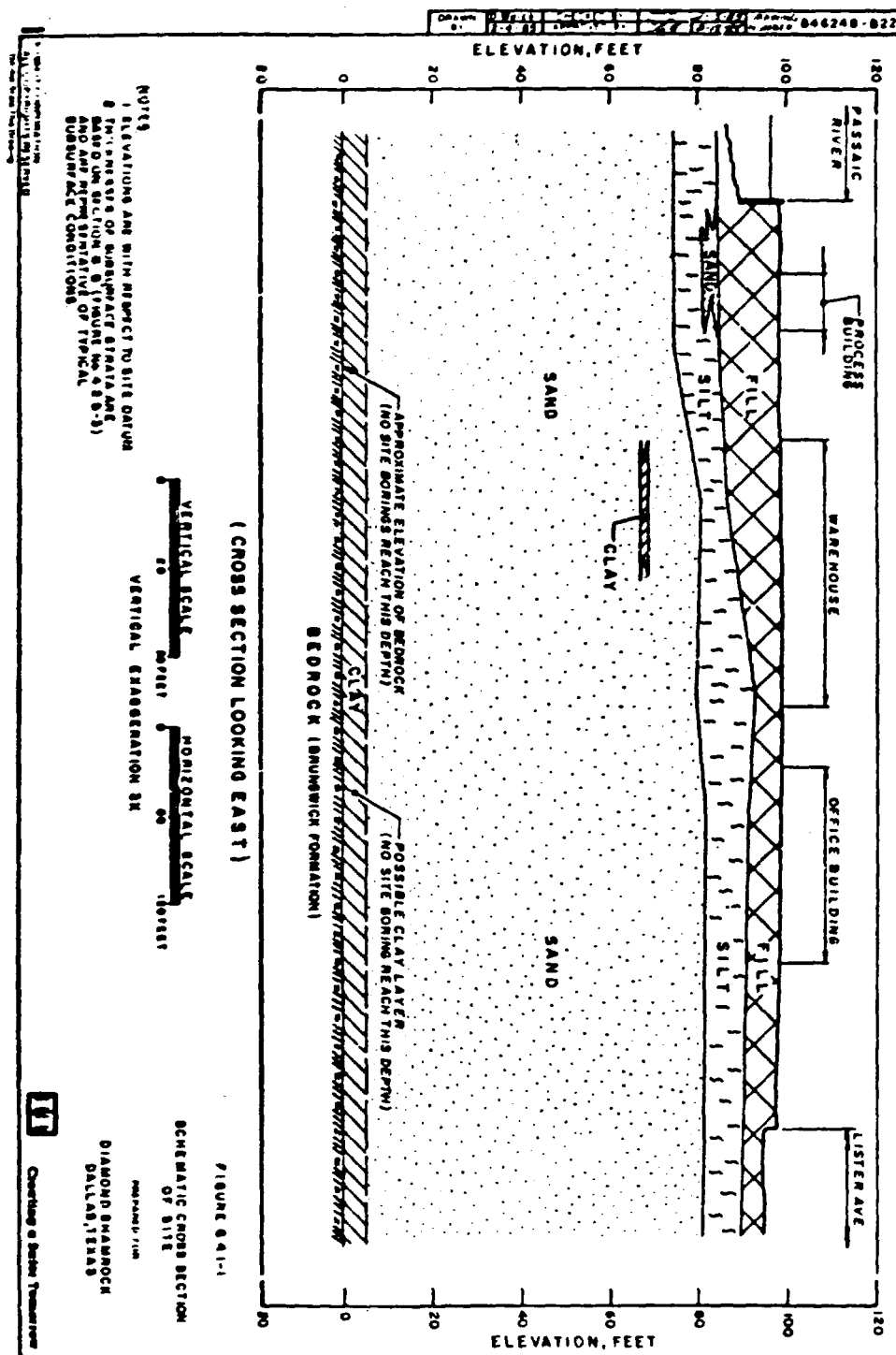
FIGURES

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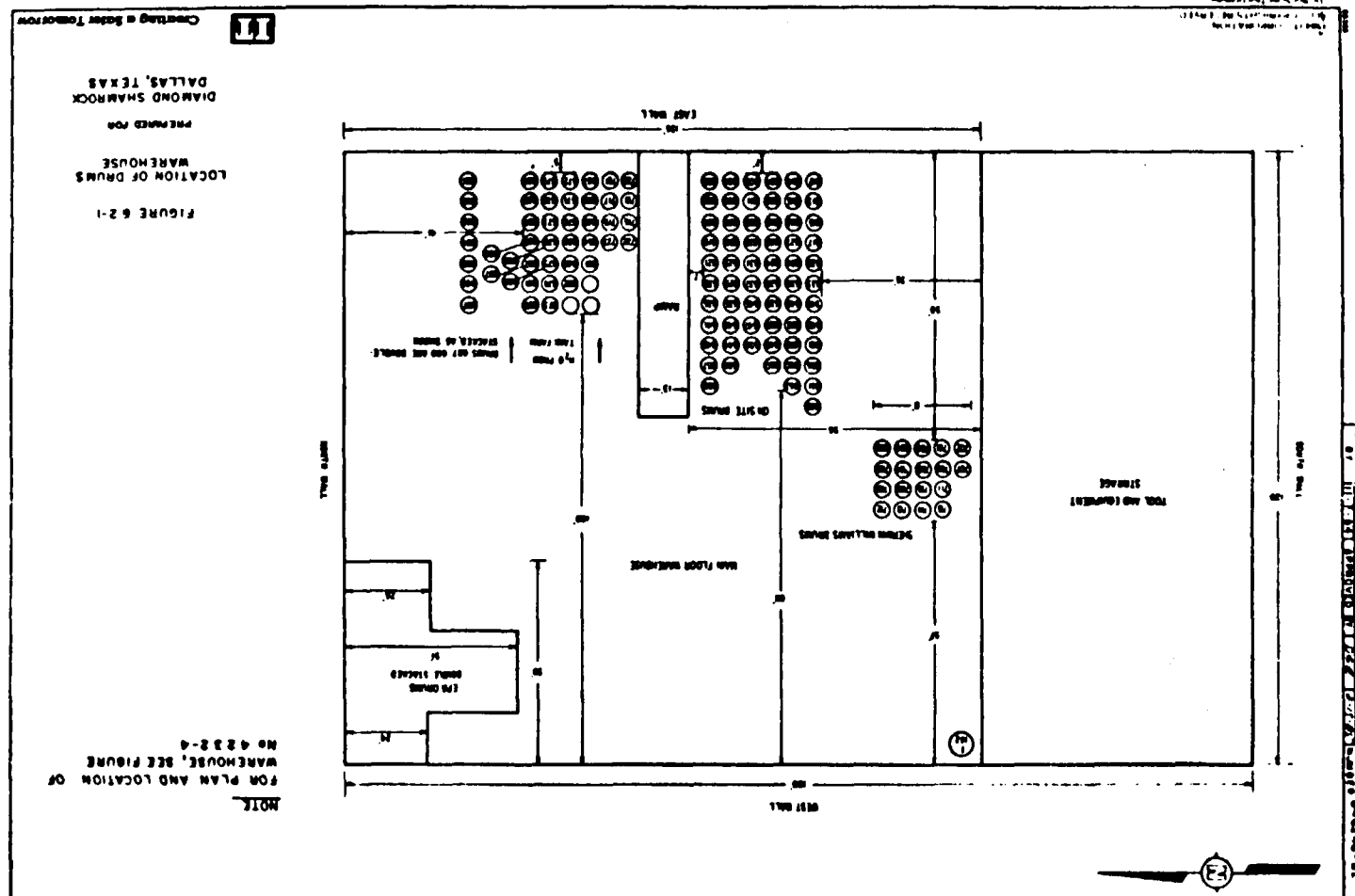
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7.0 CONCLUSIONS AND RECOMMENDATIONS

The investigation documented in this report has provided considerable data related to existing site conditions. Some conclusions regarding the extent of contamination and the need for additional data can be drawn from the results obtained thus far. Following is a brief discussion of the conclusions drawn to date and recommendations for future activities to provide the necessary input for a feasibility study.

7.1 DRUMS

Of the 570 drums on site from former plant operations, 180 have been demonstrated to have nondetectable dioxin concentrations. These drums should be segregated, externally decontaminated, and disposed of off site. The contents of the drums generated by the NJDEP, EPA, and Diamond Shamrock during off-site remedial activities and initial site stabilization should be combined with other like materials (i.e., decontamination water, tyvek clothing, etc.) currently being generated on site; the resulting empty drums could be used for continuing activities on site or decontaminated and removed from the site. The materials remaining after this consolidation and segregation will be addressed as part of the feasibility study.

7.2 BUILDINGS

With the possible exception of the office and laboratory building, the buildings on the site are highly contaminated and constructed of materials that make decontamination impractical. It is anticipated that the feasibility study will recommend demolition of these structures. Decontamination of nonporous materials, such as structural steel members, to permit their removal from the site for disposal will be evaluated as part of the feasibility study, as will the final disposal of the building rubble.

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through the silt layer into the glacio-fluvial sand aquifer or into the bedrock of the Brunswick Formation.

The subsurface hydrogeology, particularly that of the aquifers underlying the silt, needs to be further defined so that potential contaminant migration and possible remedial measures can be evaluated. This information will be an important input to the feasibility study.

7.7 RIVER SEDIMENTS

The river sediment sampling program confirmed the EPA data for the surface sediments and extended the data base to include sediments at a greater depth. The lower sediments have been shown to contain higher levels of dioxin than the shallow sediments. The lateral and vertical extent of this contamination has not been determined.

A more extensive sediment sampling program will be prepared using geostatistics as a planning tool. A hydrographic survey of the river above and below the site has already been performed. Data from this survey will be used to prepare a sediment sampling program to define the extent of river sediment contamination.

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